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<p>(21) International Application Number: PCT/US94/10264</p> <p>(22) International Filing Date: 13 September 1994 (13.09.94)</p> <p>(30) Priority Data:</p> <table border="0"> <tr> <td>08/122,230</td> <td>17 September 1993 (17.09.93)</td> <td>US</td> </tr> <tr> <td>08/122,827</td> <td>17 September 1993 (17.09.93)</td> <td>US</td> </tr> <tr> <td>08/162,827</td> <td>3 December 1993 (03.12.93)</td> <td>US</td> </tr> <tr> <td>08/172,331</td> <td>22 December 1993 (22.12.93)</td> <td>US</td> </tr> </table> <p>(71) Applicants: NOVO NORDISK A/S [DK/DK]; Novo Allé, DK-2880 Bagsværd (DK); NOVO NORDISK BIOTECH, INC. [US/US]; 1445 Drew Avenue, Davis, CA 95616-4880 (US).</p> <p>(72) Inventors: WAHLEITHNER, Jill, Angela; 1718 Tea Place, Davis, CA 95616 (US); CHRISTENSEN, Bjørn, Eggert; Dronninggaards Allé 32, DK-2840 Holte (DK); SCHNEIDER, Palle; Rydstøften 43, DK-2750 Ballerup (DK).</p> <p>(74) Agents: ZELSON, Steve, T. et al.; Novo Nordisk of North America, Inc., Suite 6400, 405 Lexington Avenue, New York, NY 10174 (US).</p>	08/122,230	17 September 1993 (17.09.93)	US	08/122,827	17 September 1993 (17.09.93)	US	08/162,827	3 December 1993 (03.12.93)	US	08/172,331	22 December 1993 (22.12.93)	US	<p>(81) Designated States: AM, AU, BB, BG, BR, BY, CA, CN, CZ, EE, FI, GE, HU, JP, KE, KG, KP, KR, KZ, LK, LR, LT, LV, MD, MG, MN, NO, NZ, PL, RO, RU, SD, SI, SK, TJ, TT, UA, UZ, VN, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG), ARIPO patent (KE, MW, SD).</p> <p>Published</p> <p><i>With international search report.</i></p> <p><i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i></p>
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<p>(54) Title: PURIFIED pH NEUTRAL RHIZOCTONIA LACCASES AND NUCLEIC ACIDS ENCODING SAME</p>													
<p>(57) Abstract</p> <p>The present invention relates to isolated nucleic acid fragments containing a sequence encoding a <i>Rhizoctonia solani</i> laccase having optimum activity at a neutral or basic pH, and the laccase proteins encoded thereby.</p>													

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PURIFIED PH NEUTRAL RHIZOCTONIA LACCASES AND NUCLEIC
ACIDS ENCODING SAME

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Related Applications

This application is a continuation-in-part of co-
pending U.S. Serial Nos. 08/122,230, 08/122,827, and
08/162,827, the contents of which are incorporated by
10 reference in their entirety.

Field of the Invention

The present invention relates to isolated nucleic acid
fragments encoding a fungal oxidoreductase enzyme and the
15 purified enzymes produced thereby. More particularly, the
invention relates to nucleic acid fragments encoding a
phenol oxidase, specifically a laccase, which functions at
a neutral pH.

20 Background of the Invention

Laccases (benzenediol:oxygen oxidoreductases) are
multi-copper containing enzymes that catalyze the oxidation
of phenolics. Laccase-mediated oxidations result in the
production of aryloxy-radical intermediates from suitable
25 phenolic substrate; the ultimate coupling of the
intermediates so produced provides a combination of dimeric,
oligomeric, and polymeric reaction products. Such reactions
are important in nature in biosynthetic pathways which lead
to the formation of melanin, alkaloids, toxins, lignins, and
30 humic acids. Laccases are produced by a wide variety of
fungi, including ascomycetes such as *Aspergillus*,
Neurospora, and *Podospora*, the deuteromycete *Botrytis*, and

basidiomycetes such as *Collybia*, *Fomes*, *Lentinus*, *Pleurotus*,
Trametes, and perfect forms of *Rhizoctonia*. Laccase
exhibits a wide range of substrate specificity, and each
different fungal laccase usually differs only quantitatively
5 from others in its ability to oxidize phenolic substrates.
Because of the substrate diversity, laccases generally have
found many potential industrial applications. Among these
are lignin modification, paper strengthening, dye transfer
inhibition in detergents, phenol polymerization, juice
10 manufacture, phenol resin production, and waste water
treatment.

Although the catalytic capabilities are similar,
laccases made by different fungal species do have different
temperature and pH optima, and these may also differ
15 depending on the specific substrate. A number of these
fungal laccases have been isolated, and the genes for
several of these have been cloned. For example, Choi et
al. (*Mol. Plant-Microbe Interactions* 5: 119-128, 1992)
describe the molecular characterization and cloning of the
20 gene encoding the laccase of the chestnut blight fungus,
Cryphonectria parasitica. Kojima et al. (*J. Biol. Chem.*
265: 15224-15230, 1990; JP 2-238885) provide a description
of two allelic forms of the laccase of the white-rot
basidiomycete *Coriolus hirsutus*. Germann and Lerch
25 (*Experientia* 41: 801, 1985; *PNAS USA* 83: 8854-8858, 1986)
have reported the cloning and partial sequencing of the
Neurospora crassa laccase gene. Saloheimo et al. (*J. Gen.*
Microbiol. 137: 1537-1544, 1985; WO 92/01046) have
disclosed a structural analysis of the laccase gene from the
30 fungus *Phlebia radiata*. However, virtually all of the
known fungal laccases function best at acidic pHs (e.g.,
between pH 3.0 and 6.0), and are typically inactive at

neutral or basic pHs. Since a number of the aforesaid potential industrial methods are preferentially conducted at neutral or basic pH, most fungal laccases perform poorly in such methods. Thus, the available fungal laccases are
5 inadequate for application in a number of important commercial methods.

An exception to this rule is the extracellular laccase produced by certain species of *Rhizoctonia*. Bollag et al. have reported a laccase with a pH optimum of about 7.0
10 produced by *Rhizoctonia praticola*. A laccase of this type would be far more useful in industrial methods requiring neutral pH than previously known laccases. However, the *R. praticola* enzyme was neither purified nor further characterized, nor, to date, has any other laccase having
15 this trait been purified or characterized. Moreover, although other laccase genes have been isolated, as described above, these have been genes encoding enzymes which function best at acidic pH. Recombinant production and commercially adequate yields of a pH neutral or basic
20 laccase have thus been unattainable due to the fact that neither the enzyme per se nor the laccase gene encoding such an enzyme has previously been isolated and/or purified and sequenced. The present invention now provides a solution to each of these problems.

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Summary of the Invention

The present invention relates to an isolated nucleic acid fragment comprising a nucleic acid sequence encoding a *Rhizoctonia* laccase which functions optimally at a pH
30 between 6.0 to 8.5. By "functioning optimally" is meant that the enzyme exhibits significant (i.e., at least about 30% of maximum, preferably at least about 50%, and most

preferably from 50% to maximum) activity within the pH range of between about 6.0-8.5, as determined by activity in one or more standard laccase assays for substrates such as the syringaldazine, ABTS, 2,6-dimethoxyphenol, or 4
5 antiaminopyrine + N-ethyl-N-sulfobutyl-m-toluidine. A preferred substrate for the laccases of the present invention is syringaldazine. In a preferred embodiment, the laccase is a *Rhizoctonia solani* laccase. The invention also relates to a substantially pure laccase encoded by the novel
10 nucleic acid sequence. By "substantially pure" is meant a laccase which is essentially (i.e., ≥90%) free of other non-laccase proteins.

In order to facilitate production of the novel laccase, the invention also provides vectors and host cells
15 comprising the claimed nucleic acid fragment, which vectors and host cells are useful in recombinant production of the laccase. The nucleic acid fragment is operably linked to transcription and translation signals capable of directing expression of the laccase protein in the host cell of
20 choice. A preferred host cell is a fungal cell, most preferably of the genus *Aspergillus*. Recombinant production of the laccase of the invention is achieved by culturing a host cell transformed or transfected with the nucleic acid fragment of the invention, or progeny thereof, under
25 conditions suitable for expression of the laccase protein, and recovering the laccase protein from the culture.

The laccases of the present invention are useful in a number of industrial processes in which oxidation of phenolics is required. These processes include lignin
30 manipulation, juice manufacture, phenol polymerization and phenol resin production. In a preferred embodiment, the

enzyme of the invention is used in a process requiring a neutral or somewhat basic pH for greatest efficiency.

Brief Description of the Figures

5 Figure 1 illustrates the nucleotide and amino acid sequence of RSlac1. Lower case letters in the nucleotide sequence indicate the position of introns.

 Figure 2 illustrates the nucleotide and amino acid sequence of RSlac2. Lower case letters in the nucleotide
10 sequence indicate the position of introns.

 Figure 3 illustrates a restriction map of the plasmid PMWR-1.

 Figure 4 illustrates the nucleotide and amino acid sequence of the translated region of RSlac3.

15 Figure 5 illustrates the syringaldazine oxidase activity of RSlac1 (90mM buffer, 20 μ M syringaldazine, 20°C).

 Figure 6 illustrates the syringaldazine oxidase activity of RSlac2 (93mM buffer, 20 μ M syringaldazine,
20 20°C).

Detailed Description of the Invention

 Certain species of the genus *Rhizoctonia* have been reported as producing laccase; therefore, an initial search focused on identifying the presence of these enzymes in
25 various *Rhizoctonia solani* isolates. Samples are cultured and the supernatants periodically analyzed for the presence of laccase by the ABTS method, described below. Laccase is observed in all the *Rhizoctonia* cultures. Harvested
 laccases are electrophoretically separated and stained with
30 ABTS. One isolate, RS22, produces a laccase with a basic pI, and is selected for further study.

The remaining studies focus on purification and characterization of the enzyme from RS22. Briefly, the fermentation broth is filtered and concentrated by UF with a membrane cut off of about 10,000. A first ion exchange chromatography step is conducted at pH 4.5 in acetate buffer, with step elution using NaCl. The eluate is then ultrafiltered and rechromatographed, and eluted with a NaCl gradient. Active fractions are pooled for further study.

The intact protein thus isolated and purified (hereinafter referred to as RSlac3) is first subjected to partial sequencing, and the N-terminal sequence obtained is as follows:

AVRNYKFDIKNVNVAPDGFQRPIVSV (SEQ. ID. NO.: 5)

The protein is further subjected to digestion with a lysine- or glutamic-acid specific protease, and additional peptides obtained from the protein have the following sequences, which can be aligned with sequences in *Coriolus hirsutus*:

Peptide 1:

SOYVDGLRGPLVIYDPDDDH (SEQ. ID. NO: 6)

Peptide 2:

GLALVFAEAPSQIRQGVQSVQPDDA (SEQ. ID. NO.: 7)

Peptide 3:

SRVBVBASTVVMLEBWYHTPAXVLE (SEQ. ID. NO. 8)

Peptide 4:

SLGPTPNYVNPXIRDVVRVGCTTVV (SEQ. ID. NO. 9)

The following peptides are also found, but do not correspond to *Coriolus* sequences

Peptide 5:

IRYVGGPAVX(N?)RSVI (SEQ. ID. NO.: 10)

Peptide 6:

ILANPA (SEQ. ID. NO.: 11)

Peptide 7:

YEAPSLPT (SEQ. ID. NO.: 12)

In the above sequences, B designates a residue which is either aspartic acid or asparagine, and X designates unidentified residues.

In order to initiate screening for a *Rhizoctonia* laccase gene, an *R. solani* genomic library is prepared. Total DNA is partially digested with restriction enzyme Sau3A, and electrophoresed in an agarose gel to isolate DNA fragments between 8 and 21 kb in size. The fractionated fragments are ligated to λ phage EMBL3 arms with BamHI ends, and the resulting phage packaged in vitro. These phage are used as a library to create a library of 170,000 plaques in *E. coli* and amplified 100-fold for future use.

In order to develop probes for isolation of the *R. solani* laccase gene, the protein sequences of five known laccases are analyzed to determine consensus sequences, and two degenerate oligonucleotides constructed based on observed consensus sequences (Choi et al. *supra*; Germann and Lerch, *supra*; Saloheimo et al, *supra*, Kojima et al, *supra*). These oligos are mixed with *R. solani* genomic DNA and a DNA fragment of 220 nucleotide fragment is amplified using a taq polymerase chain reaction(PCR). The 220-nucleotide fragment is then cloned into plasmid vector.

The PCR fragment is used as a probe to screen 25,000 plaques from the amplified genomic library. Positive clones from this screen fall into two classes that are subsequently shown, by DNA sequence analysis, to code for two different laccase genes, RSlac1 and RSlac2. The nucleotide sequence for each of these genes (SEQ ID. NOS.: 1 and 3), and the predicted amino acid sequence for each protein (SEQ. ID. NOS.: 2 and 4), are presented in, respectively, Figures 1

and 2. The homology between the two sequences is approximately 63%. Compared to known laccase sequences from *Coriolus hirsutus*, *Phlebia radiata*, *Aspergillus nidulans*, *Cryphonectria parasitica*, and *Neurospora crassa*, the RS laccases show between about 30-40% homology. Each of the two coding sequences is cloned into an expression vector operably linked to *Aspergillus oryzae* taka-amylase transcription and translation signals (See Figure 3). Each of the two laccase expression vectors is transformed into an *Aspergillus oryzae* and *Aspergillus niger* host cell, and the host cells screened for the presence of laccase.

For isolation of the RSlac3 gene, polyA RNA is purified from *R. solani* mycelia grown in the presence of anisidine. The RNA is used as a template for cDNA synthesis. The cDNA is fractionated and fragments between 1.7-3.5 kb collected, and a cDNA library created by cloning the fractionated DNA into a yeast vector. 3000 transformants from this library are screened on ABTS. After 24 hours, a single colony appears positive. The plasmid from the colony is isolated and the insert sequenced. Portions of the predicted amino acid sequence correspond with the sequences of the fragments obtained from RS 22, described *supra*. The complete nucleotide and amino acid sequences are depicted in Figure 4, and in SEQ. ID. NOS.: 13 and 14, respectively. RSlac3 shows 48% homology with RSlac1 and 50% homology with RSlac2. RSlac3 also shows 48% homology with the *Coriolus hirsutus* laccase gene.

According to the invention, a *Rhizoctonia* gene encoding a pH neutral or basic laccase can be obtained by methods described above, or any alternative methods known in the art, using the information provided herein. The gene can be expressed, in active form, using an expression

vector. A useful expression vector contains an element that permits stable integration of the vector into the host cell genome or autonomous replication of the vector in a host cell independent of the genome of the host cell, and preferably one or more phenotypic markers which permit easy selection of transformed host cells. The expression vector may also include control sequences encoding a promoter, ribosome binding site, translation initiation signal, and, optionally, a repressor gene or various activator genes. To permit the secretion of the expressed protein, nucleotides encoding a signal sequence may be inserted prior to the coding sequence of the gene. For expression under the direction of control sequences, a laccase gene to be treated according to the invention is operably linked to the control sequences in the proper reading frame. Promoter sequences that can be incorporated into plasmid vectors, and which can direct the transcription of the laccase gene, include but are not limited to the prokaryotic β -lactamase promoter (Villa-Komaroff, et al., 1978, Proc. Natl. Acad. Sci. U.S.A. 75:3727-3731) and the tac promoter (DeBoer, et al., 1983, Proc. Natl. Acad. Sci. U.S.A. 80:21-25). Further references can also be found in "Useful proteins from recombinant bacteria" in Scientific American, 1980, 242:74-94; and in Sambrook et al., Molecular Cloning, 1989.

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The expression vector carrying the DNA construct of the invention may be any vector which may conveniently be subjected to recombinant DNA procedures, and the choice of vector will typically depend on the host cell into which it is to be introduced. Thus, the vector may be an autonomously replicating vector, i.e. a vector which exists as an extrachromosomal entity, the replication of which is

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independent of chromosomal replication, e.g. a plasmid, or an extrachromosomal element, minichromosome or an artificial chromosome. Alternatively, the vector may be one which, when introduced into a host cell, is integrated into the host cell genome and replicated together with the chromosome(s) into which it has been integrated.

In the vector, the DNA sequence should be operably connected to a suitable promoter sequence. The promoter may be any DNA sequence which shows transcriptional activity in the host cell of choice and may be derived from genes encoding proteins either homologous or heterologous to the host cell. Examples of suitable promoters for directing the transcription of the DNA construct of the invention, especially in a bacterial host, are the promoter of the *lac* operon of *E.coli*, the *Streptomyces coelicolor* agarase gene *dagA* promoters, the promoters of the *Bacillus licheniformis* α -amylase gene (*amyL*), the promoters of the *Bacillus stearothermophilus* maltogenic amylase gene (*amyM*), the promoters of the *Bacillus amyloliquefaciens* α -amylase (*amyQ*), or the promoters of the *Bacillus subtilis* *xylA* and *xylB* genes. In a yeast host, a useful promoter is the *eno-1* promoter. For transcription in a fungal host, examples of useful promoters are those derived from the gene encoding *A. oryzae* TAKA amylase, *Rhizomucor miehei* aspartic proteinase, *A. niger* neutral α -amylase, *A. niger* acid stable α -amylase, *A. niger* or *A. awamsii* glucoamylase (*gluA*), *Rhizomucor miehei* lipase, *A. oryzae* alkaline protease, *A. oryzae* triose phosphate isomerase or *A. nidulans* acetamidase. Preferred are the TAKA-amylase and *gluA* promoters.

The expression vector of the invention may also comprise a suitable transcription terminator and, in eukaryotes, polyadenylation sequences operably connected to the DNA sequence encoding the laccase of the invention.

5 Termination and polyadenylation sequences may suitably be derived from the same sources as the promoter. The vector may further comprise a DNA sequence enabling the vector to replicate in the host cell in question. Examples of such sequences are the origins of replication of plasmids pUC19,
10 pACYC177, pUB110, pE194, pAMB1 and pIJ702.

The vector may also comprise a selectable marker, e.g. a gene the product of which complements a defect in the host cell, such as the *dal* genes from *B. subtilis* or *B. li-*
15 *cheniformis*, or one which confers antibiotic resistance such as ampicillin, kanamycin, chloramphenicol or tetracycline resistance. Examples of *Aspergillus* selection markers include *amdS*, *pyrG*, *argB*, *niaD* and *sc*, a marker giving rise to hygromycin resistance. Preferred for use in an
20 *Aspergillus* host cell are the *amdS* and *pyrG* markers of *A. nidulans* or *A. oryzae*. A frequently used mammalian marker is the dihydrofolate reductase (DHFR) gene. Furthermore, selection may be accomplished by co-transformation, e.g. as described in WO 91/17243.

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It is generally preferred that the expression is extracellular. The laccases of the present invention may thus comprise a preregion permitting secretion of the expressed protein into the culture medium. If desirable,
30 this preregion may be native to the laccase of the invention or substituted with a different preregion or signal sequence, conveniently accomplished by substitution of the

DNA sequences encoding the respective preregions. For example, the preregion may be derived from a glucoamylase or an amylase gene from an *Aspergillus* species, an amylase gene from a *Bacillus* species, a lipase or proteinase gene from
5 *Rhizomucor miehei*, the gene for the α -factor from *Saccharomyces cerevisiae* or the calf prochymosin gene. Particularly preferred, when the host is a fungal cell, is the preregion for *A. oryzae* TAKA amylase, *A. niger* neutral amylase, the maltogenic amylase form *Bacillus* NCIB 11837, *B.*
10 *stearothermophilus* α -amylase, or *Bacillus licheniformis* subtilisin. An effective signal sequence is the *A. oryzae* TAKA amylase signal, the *Rhizomucor miehei* aspartic proteinase signal and the *Rhizomucor miehei* lipase signal.

15 The procedures used to ligate the DNA construct of the invention, the promoter, terminator and other elements, respectively, and to insert them into suitable vectors containing the information necessary for replication, are well known to persons skilled in the art (cf., for instance,
20 Sambrook et al. Molecular Cloning, 1989).

The cell of the invention either comprising a DNA construct or an expression vector of the invention as defined above is advantageously used as a host cell in the
25 recombinant production of a enzyme of the invention. The cell may be transformed with the DNA construct of the invention, conveniently by integrating the DNA construct in the host chromosome. This integration is generally considered to be an advantage as the DNA sequence is more
30 likely to be stably maintained in the cell. Integration of the DNA constructs into the host chromosome may be performed

according to conventional methods, e.g. by homologous or heterologous recombination. Alternatively, the cell may be transformed with an expression vector as described above in connection with the different types of host cells.

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The host cell may be selected from prokaryotic cells, such as bacterial cells. Examples of suitable bacteria are gram positive bacteria such as *Bacillus subtilis*, *Bacillus licheniformis*, *Bacillus lentus*, *Bacillus brevis*, *Bacillus*
10 *stearothermophilus*, *Bacillus alkalophilus*, *Bacillus amyloliquefaciens*, *Bacillus coagulans*, *Bacillus circulans*, *Bacillus lautus*, *Bacillus megaterium*, *Bacillus thuringiensis*, or *Streptomyces lividans* or *Streptomyces murinus*, or gram negative bacteria such as *E. coli*. The
15 transformation of the bacteria may for instance be effected by protoplast transformation or by using competent cells in a manner known per se.

The host cell may also be a eukaryote, such as mammalian cells, insect cells, plant cells or preferably
20 fungal cells, including yeast and filamentous fungi. For example, useful mammalian cells include CHO or COS cells. A yeast host cell may be selected from a species of *Saccharomyces* or *Schizosaccharomyces*, e.g. *Saccharomyces cerevisiae*. Useful filamentous fungi may selected from a
25 species of *Aspergillus*, e.g. *Aspergillus oryzae* or *Aspergillus niger*. Alternatively, a strain of a *Fusarium* species, e.g. *F. oxysporum*, can be used as a host cell. Fungal cells may be transformed by a process involving protoplast formation and transformation of the protoplasts followed by regeneration of the cell wall in a manner known per
30 se. A suitable procedure for transformation of *Aspergillus* host cells is described in EP 238 023. A suitable method of

transforming *Fusarium* species is described by Malardier et al., 1989.

The present invention thus provides a method of producing a recombinant laccase of the invention, which
5 method comprises cultivating a host cell as described above under conditions conducive to the production of the enzyme and recovering the enzyme from the cells and/or culture medium. The medium used to cultivate the cells may be any conventional medium suitable for growing the host cell in
10 question and obtaining expression of the laccase of the invention. Suitable media are available from commercial suppliers or may be prepared according to published formulae (e.g. in catalogues of the American Type Culture Collection).

15 The resulting enzyme may be recovered from the medium by conventional procedures including separating the cells from the medium by centrifugation or filtration, precipitating the proteinaceous components of the supernatant or filtrate by means of a salt, e.g. ammonium sulphate, followed
20 by purification by a variety of chromatographic procedures, e.g. ion exchange chromatography, gel filtration chromatography, affinity chromatography, or the like. Preferably, the isolated protein is about 90% pure as determined by SDS-PAGE, purity being most important in food,
25 juice or detergent applications.

In a particularly preferred embodiment, the expression of laccase is achieved in a fungal host cell, such as *Aspergillus*. As described in detail in the following examples, the laccase gene is ligated into a plasmid
30 containing the *Aspergillus oryzae* TAKA α -amylase promoter, and the *Aspergillus nidulans* *amdS* selectable marker. Alternatively, the *amdS* may be on a separate plasmid and

used in co-transformation. The plasmid (or plasmids) is used to transform an *Aspergillus* species host cell, such as *A. oryzae* or *A. niger* in accordance with methods described in Yelton et al. (PNAS USA 81: 1470-1474, 1984).

5 Those skilled in the art will recognize that the invention is not limited to use of the nucleic acid fragments specifically disclosed herein, for example, in Figures 1 and 2. It will be apparent that the invention also encompasses those nucleotide sequences that encode the
10 same amino acid sequences as depicted in Figures 1, 2 and 3, but which differ from those specifically depicted nucleotide sequences by virtue of the degeneracy of the genetic code. In addition, the invention also encompasses other nucleotide fragments, and the proteins encoded thereby, which encode
15 laccase proteins having substantially the same pH optimum as those of *Rhizoctonia solani*, and which show a significant level of homology with the *Rhizoctonia solani* amino acid sequence. For example, the present data show that more than one species of *Rhizoctonia* produces a laccase with the
20 desired pH profile; it is therefore expected that other *Rhizoctonia* species also produce similar laccases and therefore, using the technology described herein, can be used as a source for genes within the scope of the claimed invention. As also shown in the present examples, not only
25 is there more than one nucleotide and amino acid sequence that encodes a laccase with the required characteristics, there is also considerable variation tolerated within the sequence while still producing a functional enzyme. Therefore, the invention also encompasses any variant
30 nucleotide sequence, and the protein encoded thereby, which protein retains at least about an 80% homology with one or the other of the amino acid sequences depicted in Figures 1,

2 and 3, and retains both the laccase and pH optimum activity of the sequences described herein. In particular, variants which retain a high level (i.e., $\geq 80\%$) of homology at highly conserved regions of the *Rhizoctonia* laccase are contemplated. Such regions are identified as residues 458-469 in RSLAC1, and 478-489 in RSLAC2; and residues 131-144 in RSLAC1 and 132-145 in RSLAC2.

Useful variants within the categories defined above include, for example, ones in which conservative amino acid substitutions have been made, which substitutions do not significantly affect the activity of the protein. By conservative substitution is meant that amino acids of the same class may be substituted by any other of that class. For example, the nonpolar aliphatic residues Ala, Val, Leu, and Ile may be interchanged, as may be the basic residues Lys and Arg, or the acidic residues Asp and Glu. Similarly, Ser and Thr are conservative substitutions for each other, as are Asn and Gln. It will be apparent to the skilled artisan that such substitutions can be made outside the regions critical to the function of the molecule and still result in an active enzyme. Retention of the desired activity can readily be determined by conducting a standard ABTS oxidation method in 0.1M sodium phosphate at pH 7.0.

The protein can be used in number of different industrial processes; although the enzyme is also functional to some extent at lower pH, the *R. solani* laccase is most beneficially used in processes that are usually conducted at a neutral or alkaline pH, since other laccases are not active in this pH range. These processes include polymerization of lignin, both Kraft and liginosulfates, in solution, in order to produce a lignin with a higher molecular weight. A neutral/alkaline laccase is a

particular advantage in that Kraft lignin is more soluble at higher pHs. Such methods are described in, for example, Jin et al., *Holzforschung* 45(6): 467-468, 1991; US Patent No. 4,432,921; EP 0 275 544; PCT/DK93/00217, 1992.

5 The laccase of the present invention can also be used for in-situ depolymerization of lignin in Kraft pulp, thereby producing a pulp with lower lignin content. This use of laccase is an improvement over the current use of chlorine for depolymerization of lignin, which leads to the
10 production of chlorinated aromatic compounds, which are an environmentally undesirable by-product of paper mills. Such uses are described in, for example, *Current opinion in Biotechnology* 3: 261-266, 1992; *J. Biotechnol.* 25: 333-339, 1992; Hiroi et al., *Svensk papperstidning* 5: 162-166, 1976.
15 Since the environment in a paper mill is typically alkaline, the present laccase is more useful for this purpose than other known laccases, which function best under acidic conditions.

Oxidation of dyes and other chromophoric compounds
20 leads to decolorization of the compounds. Laccase can be used for this purpose, which can be particularly advantageous in a situation in which a dye transfer between fabrics is undesirable, e.g., in the textile industry and in the detergent industry. Methods for dye transfer inhibition
25 and dye oxidation can be found in WO 92/01406, WO 92/18683, EP 0495836 and Calvo, *Mededelingen van de Faculteit Landbouw-wetenschappen/Rijksuniversitet Gent*.56: 1565-1567, 1991.

The present laccase can also be used for the
30 polymerization of phenolic compounds present in liquids. An example of such utility is the treatment of juices, such as apple juice, so that the laccase will accelerate a

precipitation of the phenolic compounds present in the juice, thereby producing a more stable juice. Such applications have been described in Stutz, Fruit processing 7/93, 248-252, 1993; Maier et al., Dt. Lebensmittel-
5 rindschau 86(5): 137-142, 1990; Dietrich et al., Fluss. Obst 57(2): 67-73, 1990. The invention is further illustrated by the following non-limiting examples.

EXAMPLES

1. Purification and characterization of *R. solani* laccase

- 10 Individual isolates of *R. solani* cultured on potato dextrose agar (Difco) are examined for laccase enzyme formation by transferring a small piece of agar containing vigorous growth to 100 ml CFM (24.0 g potato dextrose broth, 3.0 g yeast extract, 1.0 ml Microelement solution
15 [0.80 g KH_2PO_4 , 0.64 g $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$, 0.11 g $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$, 0.80 g $\text{MnCl}_2 \cdot 4\text{H}_2\text{O}$, 0.15 g $\text{ZnSO}_4 \cdot 7\text{H}_2\text{O}$, distilled water to 1000 ml], distilled water to 1000 ml) in a 500 ml shake flask. Incubation is at room temperature, at 200 rpm on an orbital shaker.
- 20 Samples are harvested at 50, 74, 122 and 170 hours, centrifuged and the clear supernatant analyzed for laccase with its ABTS (ABTS= 2,2'-azinobis (3 ethylbenzothiazoline-6-sulfonic acid). The analysis is carried out by adding 200 μl of 2mM ABTS in 0.1 M phosphate buffer, pH 7, and
25 observing the change in absorbance at 418 nm after 30 minutes incubation at room temperature (approximately 23-25° C). This method is modified from a peroxidase analysis method described by Pütter and Becker (Peroxidases, in: Bergmeyer, H.U. (ed.), Methods of Enzymatic Analysis, 3rd
30 ed., Vol.III, pp.286-293, 1983)

Each of the laccases harvested at 172 hours is electrophoretically separated and stained with ABTS as

chromogen. Several distinct patterns emerge; the strain RS 22 is shown to produce a laccase having a basic pI, and is chosen for further characterization.

Laccase activity is also determinable by the
5 syringaldazine method. Laccase catalyzes the oxidation of syringaldazine to tetramethoxy azo bis-methylene quinone under aerobic conditions, with a change of color from yellow to violet. 3000 μ l of 25 mM acetate buffer (containing 10mg/l cuprisulfate, 5 H₂O) at pH 5.5, 30°C, is mixed in a 1
10 cm cuvette with 225 μ l 0.28 mM syringaldazine (5mg solubilized in 25 ml ethanol and adjusted to 50 ml with demineralized water). The mixture is then mixed with 100 μ l of a laccase dilution (diluted in acetate buffer so that the increase in absorbance(AOD) is within the range of 0.1-0.6).
15 The reaction mixture is placed in a 30°C thermostated spectrophotometer and the reaction is followed at 530 nm for 10 to 70 seconds from the addition of laccase. The activity of the enzyme is calculated as AOD/minute x 0.677 x dilution factor, and is expressed as LACU.

20 For purification of the *Rhizoctonia* laccase, 2.1 liter of culture medium with a LACU activity of 0.19 LACU/ml is filtered through a 10 μ m filter and concentrated to 230 ml by ultrafiltration using a Filtron Minisette OMEGA membrane with a cutoff value of 10 kDa. The pH of the sample is 5.3
25 and the activity of the concentrated sample is determined to be 3.34 LACU/ml.

After pH adjustment to 4.5 and filtration due to slight precipitation, the sample is applied to a 40 ml S Sepharose Fast Flow column equilibrated with 20mM acetate buffer at pH
30 4.5 (buffer A). The column is washed in buffer A and eluted with buffer A containing 1 M NaCl. Active fractions are collected and pooled. This active pool is concentrated and

buffer exchanged to buffer A using an Amicon ultrafiltration unit equipped with a Diaflo YM10 membrane. This sample is rechromatographed on a 5 ml S Sepharose High Performance column using the method described above except that elution
 5 is carried out with a linear gradient over 30 column volumes from buffer A to buffer A containing 0.5 M NaCl. The fractions from this purification exhibiting highest activity are pooled. Approximately 45 mg laccase are obtained, when protein concentration is estimated by one absorption unit at
 10 A280 nm equal to 1mg/ml. The protein is >90% pure as judged by SDS-PAGE. The molecular weight estimated by SDS-PAGE is approximately 67 kDa. The specific activity of the purified protein is 1 LACU/mg. The pH profile of the purified protein, using syringaldazine as substrate is show in Table
 15 1, below.

Table 1.

pH	5	6	7	8
20 % activity	0.5	31	100	59

For sequencing of the protein, peptides are generated using wither a lysine-specific protease from *Achromobacter* (*Achromobacter* protease I) or a glutamic acid specific
 25 protease from *Bacillus licheniformes*. The peptides are purified by reverse phase HPLC employing linear gradients of 80% 2-propanol containing 0.08% aqueous TFA (solvent B) in 0.1% aqueous TFA (solvent A).

N-terminal amino acid sequence analysis of the intact
 30 protein and of purified peptides are carried out in an Applied Biosystems 473A protein sequencer according to the manufacturer's instructions. Initial partial sequencing of

the isolated protein yields the following N-terminal sequence:

AVRNYKFDIKNVNVAPDGFQRPVSV (SEQ. ID. NO.: 5)

The protein is then digested with either a lysine- or
5 glutamic-acid specific protease, and following additional peptides identified. Peptides 1-4 can be aligned with sequences in the laccase of *Coriolus hirsutus*:

Peptide 1:

SQYVDGLRGPLVIYDPDDDH (SEQ. ID. NO: 6)

10 Peptide 2:

GLALVFAEAFSQIRQGVQSVQPDDA (SEQ. ID. NO.: 7)

Peptide 3:

SRVBVBASTVVMLEBWYHTPAXVLE (SEQ. ID. NO. 8)

Peptide 4:

15 SLGPTFNYVNPXIRDVVRVGGTTVV (SEQ. ID. NO. 9)

Peptide 5:

IRYVGGPAVX(N?)RSVI (SEQ. ID. NO.: 10)

Peptide 6:

ILANPA (SEQ. ID. NO.: 11)

20 Peptide 7:

YEAPSLFT (SEQ. ID. NO.: 12)

An X in the above sequences designates an unidentified residue, and B represents a residue which is either aspartic acid or asparagine.

25

2. Isolation of *R. solani* laccase gene

A study of the known amino acid sequences of fungal laccases obtained from non-*Rhizoctonia* species (Choi et al., *supra*; German et al., *supra*; Saloheimo et al. *supra*; and
30 Kojima et al, *supra*) is conducted to determine the presence of consensus sequences among them. Two regions of high identity, IHWHGFFQ and TFWYHSH, are found near the amino

terminal third of the protein. Based on these consensus sequences and the corresponding DNA sequences, three degenerate oligonucleotides, O-lac2

[TGG/AAAGACCATA/GGTGTCTG/AGTA/G], its complement O-lac2r, and
5 O-lac3[ATCCAT/CTGGCAT/CGGG/CA/TTCTTCCAG/A], are synthesized using an Applied Biosystems 394 DNA/RNA synthesizer.

The synthesized oligos are used in a polymerase chain reaction (PCR) to screen *Rhizoctonia solani* genomic DNA for a laccase gene or fragment thereof. For amplifications of
10 genomic DNA, 0.5 µg of genomic DNA is incubated with 1µM of each primer, 200µM of dNTPs, and 1 U taq polymerase (Boehringer Mannheim) in [10 mM Tris-Cl, 1.5 mM MgCl₂, 50 mM KCl, 1 mg/ml gelatine; pH 8.3]. The reactions are incubated for 1x5 minutes at 95°C, 30x[1 minute at 95°C, 1 minute at
15 50-60°C, 1 minute at 72°C], and 1x5 minutes at 72°C. The PCR reactions amplify a DNA fragment of 220 nucleotides. The PCR product is cloned, according to manufacturer's directions, into the TA cloning vector (Invitrogen Corp.). Characterization of the PCR product by DNA sequencing of
20 individual clones distinguishes two separate laccase genes designated RSlac1 and RSlac2.

To prepare a *R. solani* genomic library, *R. solani* DNA is partially digested with restriction enzyme Sau3A, and electrophoresed through a 0.8% Sea Plaque Agarose (FMC
25 Bioproducts) in a Tris/Acetate/EDTA buffer to isolate those DNA fragments between 8.0 and 21 kb in size. The gel fractionated fragments are further purified with Beta-Agarase (New England Biolabs) according to manufacturer's instruction, and then ligated to lambda phage EMBL3 arms
30 with BamHI ends. The resulting phages are packaged in vitro using Gigapack II packaging extract (Stratagene). 25 ml of TB media+0.2% maltose and 10 MgSO₄ is inoculated into a 50 µl

aliquot of an overnight culture of *E. coli* K802 (supE, hsdR, gal, metB) and incubated at 37°C with shaking until the A600=0.5. 25 µl of a 1:10 and 1:50 dilution of the packaged phage are mixed with 250 µl of the K802 cells, and incubated
5 for 20 minutes at 37°C. To each dilution, 5 µl of melted top agar at 48°C are added. The mix is then plated onto prewarmed LB plates and incubated at 37°C for at least 12 hours. From these phage, a library of 170,000 plaques in *E. coli* K802 is created and amplified 100-fold for future
10 use.

To screen for the laccase gene, 25,000 plaques from the amplified genomic library are plated onto NZY/agarose plates for plaque lifts using conventional methods. Filters are screened using the 220 nucleotide PCR fragment randomly
15 labelled to 5×10^6 cpm/µg as a probe. Filters are hybridized in 50% formamide, 6xSSC for 16 hours at 42°C and washed with 0.5xSSC, 0.1% SDS at 65°C. Positive clones are picked and rescreened using conventional methods. The nine positive clones identified fell into two classes that by DNA sequence
20 analysis are shown to code for two different laccase genes, RSlac1 and RSlac2. The complete nucleotide sequence of each of these genes is determined using fluorescent nucleotides and an Applied Biosystems automatic DNA sequencer (Model 363A, version 1.2.0). The nucleotide and predicted amino
25 acid sequences are depicted in Figures 1 and 2.

For isolation of RSlac3, poly A RNA purified from *R. solani* mycelia grown in the presence of 1 mM anisidine is used as a template for cDNA synthesis using standard
protocols. The cDNA is fractionated by electrophoresis
30 through a 0.8% agarose gel and DNA fragments between 1.7 and 3.5 kb in size are collected. A library is then created by cloning the size-fractionated cDNA into the yeast expression

vector pYES2. 3000 yeast transformants from this library are plated initially on YNB (1.7 g yeast nitrogen base without amino acids, 5 g $(\text{NH}_4)_2\text{SO}_4$ per liter) with 2% glucose. After 4 days growth at 30°C, the resulting colonies are replica plated to YNB with 0.1% glucose, 2% galactose and 2mM ABTS [2,2'-azino-bis(3-ethylbenzthiazoline-6-sulfonic acid; Sigma # A-1888). After 24 hours of growth at 30°C a single colony has a light green halo which gradually turns a dark purple. The plasmid from this colony is isolated and the insert sequenced. The sequence of the translated portion of the RSlac3 gene and protein is shown in SEQ.ID NOS. 13 and 14, and in Figure 4.

3. Expression of laccase gene

The plasmid pmWR-1 is a pUC derived vector containing the TAKA amylase transcription regulation signals and the TAKA amylase signal sequence. This plasmid is engineered with a unique SfiI site at the signal sequence cleavage site, and a 3' adjacent NsiI site such that these two restriction enzymes can be used to introduce, in frame, a foreign protein. Using a PCR reaction (conducted as described above, but with 100 ng of the appropriate linearized plasmid DNA as a template) and mutagenized primers, an SfiI site is introduced at amino acid 12 and amino acid 14 of RSlac1 and RSlac2, respectively, such that the protein coding sequences are in frame with the TAKA signal sequence. In addition, a PCR amplification is also used to introduce a PstI site (CTGCAG) at the 3' end of RSlac1 and an NsiI site (ATGCAT) at the 3' end of RSlac2.

To prepare for transformation, cells of *Aspergillus oryzae* are cultivated in YPG (1g/l yeast extract, 0.25 g K_2PO_4 , 0.125 g/ MgSO_4 , 3.75 g glucose) at 34°C with 100-120rpm

for 16-20 hours, then collected by filtration with miracloth. Cells are washed with Mg solution (0.6M $\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$), then 2-6 g of cells are taken up in 10 ml MgP (1.2M $\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$, 10mM $\text{NaH}_2\text{PO}_4 \cdot 2\text{H}_2\text{O}$; pH 5.8). To this is
5 added 1 ml of Novozyme® 234 (120 mg/ml MgP), and the sample kept on ice for 5 minutes. One ml of BSA (12 mg/ml) is added, and the sample shaken gently at 34-37°C. Protoplasts are collected by filtration through miracloth, and overlain with 5 ml of ST (0.6 M Sorbitol, 100mM Tris; pH 7). The
10 sample is spun at 2500 rpm for 15 minutes, and a band of protoplasts collected. Two volumes of STC (1.2M Sorbitol, 10 mM tris, 10 mM $\text{CaCl}_2 \cdot 2\text{H}_2\text{O}$; pH 7.5) are added and the sample is spun at 2500 rpm for 5 minutes. The precipitate is washed twice with 5 ml of STC, and the protoplasts suspended in
15 0.5-1ml of STC.

For the transformation process, the protoplast concentration is adjusted to $1-5 \times 10^7/\text{ml}$. To 100 μl of protoplast solution is added a maximum of 10 μl of DNA solution (5-10 μg of supercoiled DNA) and 0.2 ml of PEG
20 (60% PEG4000, 10mM Tris, 10mM $\text{CaCl}_2 \cdot \text{H}_2\text{O}$; pH 7.5), and the combination is mixed well. The sample is kept at room temperature for 25 minutes; then to it is added first 0.2 ml PEG, with mixing, the 0.85 ml PEG with mixing. The mixture is kept at room temperature for 20 minutes, then spun at
25 4000 rpm for 15 minutes. The precipitate is washed with 2 ml of STC by spinning at 2500 rpm for 10 minutes. The protoplasts are resuspended in 0.2-0.5 ml of STC, and then spread on COVE plates. COVE medium (pH 7) contains 342.3 g/l sucrose, 25 g/l agar and a salt solution comprising 26 g/l
30 KCl, 26 g/l $\text{MgSO}_4 \cdot \text{H}_2\text{O}$, 76 g/l KH_2PO_4 , and 50 ml/l of trace metals; the trace metals are 40 mg/l $\text{NaB}_4\text{O}_7 \cdot 10\text{H}_2\text{O}$, 400 mg/l

CuSO₄·5H₂O, 1200mg/l FeSO₄·7H₂O, 700mg/l MnSO₄·H₂O, 800mg/l Na₂MoO₄·2H₂O, 10 g/l ZnSO₄·7H₂O). After autoclaving, 10 ml/l of 1M filtrated acetamide and 5-10 ml of 3M CsCl are added to the solution. Transformants are selected by growth cells
5 on COVE medium which contains acetamide as the carbon source.

The confirmation of laccase production in the samples is determined by the ABTS oxidation method as described above on Cove medium with 2 mM ABTS, at pH 5 and 7.3. Both
10 RSlac1 and RSlac2 express laccase activity at pH 5 and pH 7, in contrast with a control laccase which shows substantially no activity at pH 7.3.

The products of the expression of each of RSlac1 and RSlac2 are tested for oxidase activity at various pHs using
15 syringaldazine as the substrate. The assay is conducted substantially as described above for the assay of the native protein, over pH range of 4-9. As shown in Figures 5 and 6, both laccases are active at pHs over pH 5, and RSlac1 has particularly good activity at pHs over 6. The pattern of
20 activity is generally comparable to that observed for the RSlac3 laccase isolated from RS 22 (see Table 1 above), with RSlac1 exhibiting the broadest range of activity.

Deposit of Biological Materials

The following biological materials have been deposited
25 under the terms of the Budapest Treaty in the International Mycological Institute, Genetic Resource Reference Collection, located at Bakeham Lane, Egham, Surrey TW20 9TY and given the following accession number.

30 <u>Deposit</u>	<u>Accession Number</u>
<i>Rhizoctonia solani</i> RS22	IMI CC 358730

The following biological materials have been deposited under the terms of the Budapest Treaty with the Agricultural Research Service Patent Culture Collection, Northern Regional Research Center, 1815 University Street, Peoria, Illinois, 61604 and given the following accession numbers.

<u>Deposit</u>	<u>Accession Number</u>
5 E. coli containing RSlac1 fused to an α -amylase signal sequence (EMCC 00844)	NRRL B-21141
10 E. coli containing RSlac2 with an SfiI site insert (EMCC 00845)	NRRL B-21142
15 E. coli containing RSlac3 (EMCC 0088)	NRRL B-21156

SEQUENCE LISTING

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(iii) NUMBER OF SEQUENCES: 14

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(v) COMPUTER READABLE FORM:

(A) MEDIUM TYPE: Floppy disk
(B) COMPUTER: IBM PC compatible
(C) OPERATING SYSTEM: PC-DOS/MS-DOS
(D) SOFTWARE: PatentIn Release #1.0, Version #1.25

(vi) CURRENT APPLICATION DATA:

(A) APPLICATION NUMBER: to be assigned
(B) FILING DATE: 13-SEP-1994

(vii) PRIOR APPLICATION DATA:

(A) APPLICATION NUMBER: US 08/172,331
(B) FILING DATE: 22-DEC-1993

(vii) PRIOR APPLICATION DATA:

(A) APPLICATION NUMBER: US 08/122,230
(B) FILING DATE: 17-SEP-1993

(vii) PRIOR APPLICATION DATA:

(A) APPLICATION NUMBER: US 08/122,827
(B) FILING DATE: 17-SEP-1993

(vii) PRIOR APPLICATION DATA:

(A) APPLICATION NUMBER: US 08/162,827
(B) FILING DATE: 03-DEC-1993

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(2) INFORMATION FOR SEQ ID NO:1:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 2838 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (vi) ORIGINAL SOURCE:
 - (A) ORGANISM: Rhizoctonia laccase
- (ix) FEATURE:
 - (A) NAME/KEY: intron
 - (B) LOCATION: 302..351
- (ix) FEATURE:
 - (A) NAME/KEY: intron
 - (B) LOCATION: 463..512
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 - (A) NAME/KEY: intron
 - (B) LOCATION: 1827..1880
- (ix) FEATURE:
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- (ix) FEATURE:

(A) NAME/KEY: intron
(B) LOCATION: 2438..2498

(ix) FEATURE:

(A) NAME/KEY: CDS
(B) LOCATION: join(170..301, 352..462, 513..575, 634..759, 819
..821, 878..1000, 1055..1315, 1373..1696, 1755
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..2437, 2499..2521)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

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Arg Thr Thr Phe Leu Val Ser Val Ser Leu Phe Val Ser Ala Val Leu
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CCT GAC GGT GTT AAG CGT AAT GCG ACT TTG GTACGCACTC CTTGTAATCC      321
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Lys Leu Thr Asn Pro Glu Met Tyr Arg Thr Thr Ser Ile
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Val Thr Val Ile Ala Ala Asp Gly Val Ser Thr Lys Pro Tyr Gln Val	
245 250 255	
GAT GCG TTT GAT ATT CTA GCA GGA CAG CGC ATA GAT TGC GTC	1315
Asp Ala Phe Asp Ile Leu Ala Gly Gln Arg Ile Asp Cys Val	
260 265 270	
GTAAGTGTCG TCCGAACCCA CATCTGAGCT CAAGTGTTGA TACATGCCCG CTTATAG	1372
GTG GAG GCG AAC CAA GAA CCC GAC ACA TAC TGG ATC AAC GCA CCG CTG	1420
Val Glu Ala Asn Gln Glu Pro Asp Thr Tyr Trp Ile Asn Ala Pro Leu	
275 280 285	
ACC AAC GTG CCC AAC AAG ACC GCT CAG GCT CTC CTC GTT TAT GAG GAG	1468
Thr Asn Val Pro Asn Lys Thr Ala Gln Ala Leu Leu Val Tyr Glu Glu	
290 295 300 305	
GAT CGT CCG CCG TAC CAC CCT CCA AAG GGC CCG TAT CCC AAC TGG AGC	1516
Asp Arg Arg Pro Tyr His Pro Pro Lys Gly Pro Tyr Arg Lys Trp Ser	
310 315 320	
GTC TCT GAG GCG ATC ATC AAG TAC TGG AAT CAC AAG CAC AAG CAC GGA	1564
Val Ser Glu Ala Ile Ile Lys Tyr Trp Asn His Lys His Lys His Gly	
325 330 335	
CGT GGT TTG CTG TCT GGA CAT GGA GGT CTC AAG GCT CGG ATG ATC GAG	1612
Arg Gly Leu Leu Ser Gly His Gly Gly Leu Lys Ala Arg Met Ile Glu	
340 345 350	
GGT AGC CAT CAT CTG CAT TCG CGC AGC GTC GTT AAG CGC CAG AAT GAG	1660

Gly	Ser	His	His	Leu	His	Ser	Arg	Ser	Val	Val	Lys	Arg	Gln	Asn	Glu		
355						360					365						
ACC	ACC	ACT	GTT	GTA	ATG	GAC	GAG	AGC	AAG	CTC	GTT	GTAACTACCA				1706	
Thr	Thr	Thr	Val	Val	Met	Asp	Glu	Ser	Lys	Leu	Val						
370					375					380							
TATTTAAAAG	TTGGTTGGGT	TTCGAATACT	TATTTCAACT	TTTCTTAG	CCA	CTG	GAA									1763	
															Pro	Leu	Glu
TAC	CCC	GGC	GCT	GCA	TGC	GGG	TCT	AAA	CCT	GCT	GAC	CTC	GTC	TTG	GAT		1811
Tyr	Pro	Gly	Ala	Ala	Cys	Gly	Ser	Lys	Pro	Ala	Asp	Leu	Val	Leu	Asp		
385					390					395					400		
CTC	ACT	TTT	GGT	TTG	GTATGTAGCC	AAATCGCCCA	TATACAGGAT	ACTGAATATT									1866
Leu	Thr	Phe	Gly	Leu													
				405													
GTTCGTCCGT	GTAG	AAC	TTT	GCT	ACC	GGG	CAC	TGG	ATG	ATC	AAC	GGT	ATC				1916
		Asn	Phe	Ala	Thr	Gly	His	Trp	Met	Ile	Asn	Gly	Ile				
						410						415					
CCA	TAC	GAG	TCT	CCC	AAA	ATC	CCC	ACA	TTG	CTC	AAG	ATC	CTC	ACT	GAT		1964
Pro	Tyr	Glu	Ser	Pro	Lys	Ile	Pro	Thr	Leu	Leu	Lys	Ile	Leu	Thr	Asp		
		420					425					430					
GAG	GAC	GGG	GTT	ACC	GAG	TCT	GAC	TTC	GTATGTCC	TTTTCCGTAT							2011
Glu	Asp	Gly	Val	Thr	Glu	Ser	Asp	Phe									
		435					440										
CTTCGTATGC	GTGCACTGAC	TCGTGCTGGT	GGGAATTTAG	ACC	AAG	GAG	GAG	CAC									2066
									Thr	Lys	Glu	Glu	His				
											445						
ACA	GTC	ATA	CTC	CCG	AAG	AAC	AAA	TGC	ATC	GAA	TTC	AAC	ATC	AAG	GGG		2114
Thr	Val	Ile	Leu	Pro	Lys	Asn	Lys	Cys	Ile	Glu	Phe	Asn	Ile	Lys	Gly		
		450					455					460					
AAC	TCG	GGT	ATT	CCC	ATT	ACG	CAC	CCC	GTA	CAT	CTT	CAC	GGT				2156
Asn	Ser	Gly	Ile	Pro	Ile	Thr	His	Pro	Val	His	Leu	His	Gly				
		465					470				475						
GTAAGTGCAT	ATCGGATGGT	TTACGATACT	AAGGCTCATC	AACTTTTTAG	CAC	ACT											2212
															His	Thr	
TGG	GAT	GTC	GTA	CAA	TTT	GGC	AAC	AAC	CCA	CCC	AAT	TAT	GTC	AAT	CCT		2260
Trp	Asp	Val	Val	Gln	Phe	Gly	Asn	Asn	Pro	Pro	Asn	Tyr	Val	Asn	Pro		
480					485					490					495		
CCC	CGT	AGG	GAC	GTG	GTT	GGC	TCT	ACA	GAT	GCG	GGT	GTG	AGG	ATT	CAG		2308
Pro	Arg	Arg	Asp	Val	Val	Gly	Ser	Thr	Asp	Ala	Gly	Val	Arg	Ile	Gln		
				500						505				510			
TTC	AAG	ACC	GAC	AAT	CCA	GGA	CCG	TGG	PTC	CTG	CAC	TGC	GTGCGTCGGT				2357
Phe	Lys	Thr	Asp	Asn	Pro	Gly	Pro	Trp	Phe	Leu	His	Cys					
			515					520									
CCCCATCGTC	CGTTATGGTT	TTTCTAATAC	GTCCCATTTCT	ATTTTAG	CAT	ATT	GAC										2413
															His	Ile	Asp
															525		
TGG	CAT	CTT	GAG	GAG	GGT	TTC	GCA	GTGAGTACTG	AGACCTAAGT	GCTACTCGGC							2467
Trp	His	Leu	Glu	Glu	Gly	Phe	Ala										
		530					535										

TCATTACTGA TTACCGCATG TATGCGTCTA G ATG GTG TTT GCT GAA GCG CCC 2519
Met Val Phe Ala Glu Ala Pro
540

GAA GCC GTC AAG GGA GGT CCA AAG AGC GTG GCC GTG GAC TCT CAG TGG 2567
Glu Ala Val Lys Gly Gly Pro Lys Ser Val Ala Val Asp Ser Gln Trp
545 550 555

GAA GGG CTG TGT GGC AAG TAC GAC AAC TGG CTA AAA TCA AAT CCG GGC 2615
Glu Gly Leu Cys Gly Lys Tyr Asp Asn Trp Leu Lys Ser Asn Pro Gly
560 565 570

CAG CTG TAGGCGTATC GCAGCCACAT TGGTGATGAT TGAAAGTTGC ATCTTGTCTCC 2671
Gln Leu
575

TATAACCGGC TCTTATATAC GGGTGTCTCC CAGTAAAGTC GTAGCCCAAT TTCAGCCGAG 2731

ACAGATATTT AGTGGACTCT TACTCTGTG TCCCATTGAC GCACATCGTT GCATCAAACC 2791

TGCTTTTAT CGTCCCTCTT TGTAAATTTGT GTTGCTGTAA TGTATCG 2838

(2) INFORMATION FOR SEQ ID NO:2:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 576 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

Met Ala Arg Thr Thr Phe Leu Val Ser Val Ser Leu Phe Val Ser Ala
1 5 10 15

Val Leu Ala Arg Thr Val Glu Tyr Gly Leu Lys Ile Ser Asp Gly Glu
20 25 30

Ile Ala Pro Asp Gly Val Lys Arg Asn Ala Thr Leu Val Asn Gly Gly
35 40 45

Tyr Pro Gly Pro Leu Ile Phe Ala Asn Lys Gly Asp Thr Leu Lys Val
50 55 60

Lys Val Gln Asn Lys Leu Thr Asn Pro Glu Met Tyr Arg Thr Thr Ser
65 70 75 80

Ile His Trp His Gly Leu Leu Gln His Arg Asn Ala Asp Asp Asp Gly
85 90 95

Pro Ser Phe Val Thr Gln Cys Pro Ile Val Pro Arg Glu Ser Tyr Thr
100 105 110

Tyr Thr Ile Pro Leu Asp Asp Gln Thr Gly Thr Tyr Trp Tyr His Ser
115 120 125

His Leu Ser Ser Gln Tyr Val Asp Gly Leu Arg Gly Pro Leu Val Ile
130 135 140

Tyr Asp Pro Lys Asp Pro His Arg Arg Leu Tyr Asp Val Asp Asp Glu
145 150 155 160

Lys Thr Val Leu Ile Ile Gly Asp Trp Tyr His Glu Ser Ser Lys Ala
165 170 175

Ile Leu Ala Ser Gly Asn Ile Thr Arg Gln Arg Pro Val Ser Ala Thr
 180 185 190
 Ile Asn Gly Lys Gly Arg Phe Asp Pro Asp Asn Thr Pro Ala Asn Pro
 195 200 205
 Asp Thr Leu Tyr Thr Leu Lys Val Lys Arg Gly Lys Arg Tyr Arg Leu
 210 215 220
 Arg Val Ile Asn Ser Ser Glu Ile Ala Ser Phe Arg Phe Ser Val Glu
 225 230 235 240
 Gly His Lys Val Thr Val Ile Ala Ala Asp Gly Val Ser Thr Lys Pro
 245 250 255
 Tyr Gln Val Asp Ala Phe Asp Ile Leu Ala Gly Gln Arg Ile Asp Cys
 260 265 270
 Val Val Glu Ala Asn Gln Glu Pro Asp Thr Tyr Trp Ile Asn Ala Pro
 275 280 285
 Leu Thr Asn Val Pro Asn Lys Thr Ala Gln Ala Leu Leu Val Tyr Glu
 290 295 300
 Glu Asp Arg Arg Pro Tyr His Pro Pro Lys Gly Pro Tyr Arg Lys Trp
 305 310 315 320
 Ser Val Ser Glu Ala Ile Ile Lys Tyr Trp Asn His Lys His Lys His
 325 330 335
 Gly Arg Gly Leu Leu Ser Gly His Gly Gly Leu Lys Ala Arg Met Ile
 340 345 350
 Glu Gly Ser His His Leu His Ser Arg Ser Val Val Lys Arg Gln Asn
 355 360 365
 Glu Thr Thr Thr Val Val Met Asp Glu Ser Lys Leu Val Pro Leu Glu
 370 375 380
 Tyr Pro Gly Ala Ala Cys Gly Ser Lys Pro Ala Asp Leu Val Leu Asp
 385 390 395 400
 Leu Thr Phe Gly Leu Asn Phe Ala Thr Gly His Trp Met Ile Asn Gly
 405 410 415
 Ile Pro Tyr Glu Ser Pro Lys Ile Pro Thr Leu Leu Lys Ile Leu Thr
 420 425 430
 Asp Glu Asp Gly Val Thr Glu Ser Asp Phe Thr Lys Glu Glu His Thr
 435 440 445
 Val Ile Leu Pro Lys Asn Lys Cys Ile Glu Phe Asn Ile Lys Gly Asn
 450 455 460
 Ser Gly Ile Pro Ile Thr His Pro Val His Leu His Gly His Thr Trp
 465 470 475 480
 Asp Val Val Gln Phe Gly Asn Asn Pro Pro Asn Tyr Val Asn Pro Pro
 485 490 495
 Arg Arg Asp Val Val Gly Ser Thr Asp Ala Gly Val Arg Ile Gln Phe
 500 505 510
 Lys Thr Asp Asn Pro Gly Pro Trp Phe Leu His Cys His Ile Asp Trp
 515 520 525
 His Leu Glu Glu Gly Phe Ala Met Val Phe Ala Glu Ala Pro Glu Ala

530		535		540
Val Lys Gly Gly Pro Lys Ser Val Ala Val Asp Ser Gln Trp Glu Gly				
545		550		555
				560
Leu Cys Gly Lys Tyr Asp Asn Trp Leu Lys Ser Asn Pro Gly Gln Leu				
	565		570	575

(2) INFORMATION FOR SEQ ID NO:3:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 3117 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(vi) ORIGINAL SOURCE:

- (A) ORGANISM: Rhizoctonia laccase

(ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: join(393..524, 577..687, 737..799, 860..985, 1043..1045, 1097..1219, 1269..1538, 1601..1996, 2047..2118, 2174..2284, 2338..2439, 2495..2635, 2693..2725, 2786..2899)

(ix) FEATURE:

- (A) NAME/KEY: intron
- (B) LOCATION: 525..576

(ix) FEATURE:

- (A) NAME/KEY: intron
- (B) LOCATION: 688..736

(ix) FEATURE:

- (A) NAME/KEY: intron
- (B) LOCATION: 800..859

(ix) FEATURE:

- (A) NAME/KEY: intron
- (B) LOCATION: 986..1042

(ix) FEATURE:

- (A) NAME/KEY: intron
- (B) LOCATION: 1220..1268

(ix) FEATURE:

- (A) NAME/KEY: intron
- (B) LOCATION: 1539..1600

(ix) FEATURE:

- (A) NAME/KEY: intron
- (B) LOCATION: 1823..1936

(ix) FEATURE:

- (A) NAME/KEY: intron
- (B) LOCATION: 1973..2046

(ix) FEATURE:

- (A) NAME/KEY: intron
- (B) LOCATION: 2119..2173

(ix) FEATURE:

- (A) NAME/KEY: intron

(B) LOCATION: 2285..2337

(ix) FEATURE:

(A) NAME/KEY: intron

(B) LOCATION: 2440..2494

(ix) FEATURE:

(A) NAME/KEY: intron

(B) LOCATION: 2636..2692

(ix) FEATURE:

(A) NAME/KEY: intron

(B) LOCATION: 1046..1096

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

GAGTGATCCG CCAGAGTTCA GCGGATAAG TTCCTAAATA GTCATTGCGC TATTCGTGTA	60
CCTCAGCATA CTGACGACAT ACCGCCAGAT CGCCCTCGGT TCGGGCGTGG CATACGTTCC	120
CAAGGGCACC TCACGGAGCA AACTCTAAAA AGCTTCGGCA TGGATTGCAT TTTGTATTGT	180
AAACAAGTTA CGAGAAAAAC AATAGATCAG TTTTTCGCGA ATCGGATGGC TTGAAACGGA	240
AGTACCGATG GCCGATCCGA GTCGAATGAA TTAACGCATC TGAAACGGGA CCCTGAGTCG	300
AGGCACCCGC CGGCCCTGGC CGTATAAGTC ACTTGTGCGC AACTAGCACT TTTTCATTCC	360
CCCTTTTCTT CTTCCTCGTC TTCTTCTTCT CT ATG GCT CGG TCG ACT ACT TCA	413
Met Ala Arg Ser Thr Thr Ser	
1 5	
CTC TTT GCA CTG TCT CTC GTT GCT TCA GCG TTT GCT CGA GTC GTT GAC	461
Leu Phe Ala Leu Ser Leu Val Ala Ser Ala Phe Ala Arg Val Val Asp	
10 15 20	
TAT GGG TTT GAT GTG GCT AAT GGG GCA GTT GCT CCG GAT GCT GTA ACA	509
Tyr Gly Phe Asp Val Ala Asn Gly Ala Val Ala Pro Asp Gly Val Thr	
25 30 35	
AGG AAC GCG GTT CTC GTGAGTTAGC TGTAAGATGG TGTATATGCT GGTTCCTAA	564
Arg Asn Ala Val Leu	
40	
CGGGAATGTC AG GTC AAT GGT CGC TTC COT GGT CCA TTG ATC ACC GCC	612
Val Asn Gly Arg Phe Pro Gly Pro Leu Ile Thr Ala	
45 50 55	
AAC AAG GGG GAT ACA CTT AAA ATC ACC GTG CGG AAT AAA CTC TCC GAT	660
Asn Lys Gly Asp Thr Leu Lys Ile Thr Val Arg Asn Lys Leu Ser Asp	
60 65 70	
CCA ACT ATG CGA AGG AGC ACG ACC ATC GTTAGTACTT CCCCTCATCT	767
Pro Thr Met Arg Arg Ser Thr Thr Ile	
75 80	
GTCTTGAAAC TTTCTCATCT TTTTGAAG CAC TCG CAC GGT CTC CTC CAA CAC	760
His Trp His Gly Leu Leu Gln His	
85	
AGG ACG GCA GAA GAA GAT GGC CCG GCC TTT GTA ACC CAG GTATGCCCTTA	809
Arg Thr Ala Glu Glu Asp Gly Pro Ala Phe Val Thr Gln	
90 95 100	
TCCTATCGCT GCTCTGTCCC CGCGTCCTTC CCTGACTCGG GCGATTCTAG TGC CCG	865
Cys Pro	

ATT CCT CCG CAA GAA TCG TAC ACC TAT ACG ATG CCG CTC GGC GAA CAG Ile Pro Pro Gln Glu Ser Tyr Thr Tyr Thr Met Pro Leu Gly Glu Gln 105 110 115 120	513
ACC GGC ACG TAT TGG TAC CAC AGC CAC TTG AGC TCC CAG TAT GTG GAC Thr Gly Thr Tyr Trp Tyr His Ser His Leu Ser Ser Gln Tyr Val Asp 125 130 135	961
GGG TTG CGT GGG CCC ATC GTT ATT GTAAGTCTTC ATTTAACCTT ATTCTTGGTT Gly Leu Arg Gly Pro Ile Val Ile 140	1015
ATGGCTGATT GTGACGTCGT GGTAGT ATG TTCGTGGCTT CCACAAGAAG Met 145	1065
TCAGCAGCCC TTGAAGCTAA CTTTATTCCA G GAC CCC CAC GAC CCG TAC AGA Asp Pro His Asp Pro Tyr Arg 150	1117
AAC TAC TAT GAT GTC GAC GAC GAG CGT ACG GTC TTT ACT TTA GCA GAC Asn Tyr Tyr Asp Val Asp Asp Glu Arg Thr Val Phe Thr Leu Ala Asp 155 160 165	1165
TGG TAC CAC ACG CCG TCG GAG GCT ATC ATT GCC ACC CAC GAT GTC TTG Trp Tyr His Thr Pro Ser Glu Ala Ile Ile Ala Thr His Asp Val Leu 170 175 180	1213
AAA ACG GTACGCGTTA ATCCTTCTAG CTTTCTTTCC TTGGGTCACT TTCTATCAG Lys Thr 185	1268
ATC CCC GAC TCG GGT ACG ATC AAC GGC AAA GGC AAA TAC GAT CCT GCT Ile Pro Asp Ser Gly Thr Ile Asn Gly Lys Gly Lys Tyr Asp Pro Ala 190 195 200	1316
TCG GCT AAC ACC AAC AAC ACG ACA CTC GAG AAC CTC TAC ACT CTC AAA Ser Ala Asn Thr Asn Asn Thr Thr Leu Glu Asn Leu Tyr Thr Leu Lys 205 210 215	1364
GTC AAA CGC GGC AAG CGG TAT CGC CTG AGG ATT ATC AAC GCC TCG GCC Val Lys Arg Gly Lys Arg Tyr Arg Leu Arg Ile Ile Asn Ala Ser Ala 220 225 230	1412
ATC GCT TCG TTC CGG TTC GGC GTG CAG GGC CAC AAG TGC ACG ATC ATC Ile Ala Ser Phe Arg Phe Gly Val Gln Gly His Lys Cys Thr Ile Ile 235 240 245 250	1460
GAG GCT GAT GGC GTC CTC ACC AAA CCG ATC GAG GTC GAT GCG TTT GAT Glu Ala Asp Gly Val Leu Thr Lys Pro Ile Glu Val Asp Ala Phe Asp 255 260 265	1508
ATT CTA GCA GGC CAG AGG TAT AGC TGC ATC GTAAGTCTAC CTATGCCTTG Ile Leu Ala Gly Gln Arg Tyr Ser Cys Ile 270 275	1558
TTGTGGAGAT AAGAACCTGA CTGAATGTAT GCGCTCCAAT AG TTG AAG GCC GAC Leu Lys Ala Asp 280	1612
CAA GAT CCT GAT TCC TAC TGG ATA AAT GCG CCA ATC ACA AAC GTT CTC Gln Asp Pro Asp Ser Tyr Trp Ile Asn Ala Pro Ile Thr Asn Val Leu 285 290 295	1660
AAC ACC AAC GTC CAG GCA TTG CTA GTG TAT GAA GAT GAC AAG CGT CCT	1708

Asn	Thr	Asn	Val	Gln	Ala	Leu	Leu	Val	Tyr	Glu	Asp	Asp	Lys	Arg	Pro		
			300					305					310				
ACT	CAC	TAC	CCC	TGG	AAG	CCG	TTT	TTG	ACA	TGG	AAG	ATA	TCA	AAT	GAA	1756	
Thr	His	Tyr	Pro	Trp	Lys	Pro	Phe	Leu	Thr	Trp	Lys	Ile	Ser	Asn	Glu		
		315					320					325					
ATC	ATT	CAG	TAC	TGG	CAG	CAC	AAG	CAC	GGG	TGG	CAC	GGT	CAC	AAG	GGA	1804	
Ile	Ile	Gln	Tyr	Trp	Gln	His	Lys	His	Gly	Ser	His	Gly	His	Lys	Gly		
	330					335					340						
AAG	GGG	CAT	CAT	CAT	AAA	GTC	CGG	GCC	ATT	GGA	GGT	GTA	TCC	GGG	TTG	1852	
Lys	Gly	His	His	His	Lys	Val	Arg	Ala	Ile	Gly	Gly	Val	Ser	Gly	Leu		
345					350				355						360		
AGC	TCC	AGG	GTT	AAG	AGC	CGG	GCG	AGT	GAC	CTA	TGG	AAG	AAG	GCT	GTC	1900	
Ser	Ser	Arg	Val	Lys	Ser	Arg	Ala	Ser	Asp	Leu	Ser	Lys	Lys	Ala	Val		
			365					370						375			
GAG	TTG	GCT	GCT	GCA	CTC	GTT	GCG	GGT	GAG	GCC	GAG	TTG	GAC	AAG	AGG	1948	
Glu	Leu	Ala	Ala	Ala	Leu	Val	Ala	Gly	Glu	Ala	Glu	Leu	Asp	Lys	Arg		
		380					385					390					
CAG	AAT	GAG	GAT	AAT	TCG	ACT	ATT	GTA	TTG	GAT	GAG	ACC	AAG	CTT	ATT	1996	
Gln	Asn	Glu	Asp	Asn	Ser	Thr	Ile	Val	Leu	Asp	Glu	Thr	Lys	Leu	Ile		
	395					400					405						
GTAAGTCCCT	TAATTTTTTTT	CGGTGTCACG	GAAGCTAACC	CGCGTAATAG	CCG	TTG										2052	
						Pro	Leu								410		
GTT	CAA	CCT	GCT	GCA	CCG	GGC	GGC	TCC	AGA	CCA	GCT	GAC	GTC	GTG	GTC	2100	
Val	Gln	Pro	Gly	Ala	Pro	Gly	Gly	Ser	Arg	Pro	Ala	Asp	Val	Val	Val		
			415					420						425			
CCT	CTG	GAC	TTT	GGC	CTC	GTA	GTG	GCT	TCT	GTT	TATT	CGT	CCG	GAAT		2148	
Pro	Leu	Asp	Phe	Gly	Leu												
		430															
GCAAACTGAT	TTGGGTGGGC	TATAG	AAC	TTT	GCC	AAC	GGA	CTG	TGG	ACG	ATA					2200	
			Asn	Phe	Ala	Asn	Gly	Leu	Trp	Thr	Ile					440	
					435												
AAC	AAT	GTC	TCC	TAC	TCC	CCT	CCG	GAT	GTC	CCT	ACT	CTC	CTC	AAG	ATC	2248	
Asn	Asn	Val	Ser	Tyr	Ser	Pro	Pro	Asp	Val	Pro	Thr	Leu	Leu	Lys	Ile		
		445					450					455					
TTG	ACC	GAC	AAA	GAC	AAA	GTC	GAC	GCT	TCT	GAC	TTC	GTA	GTT	CTC		2294	
Leu	Thr	Asp	Lys	Asp	Lys	Val	Asp	Ala	Ser	Asp	Phe						
		460				465											
CTTCTTCTTT	TCAAAC TAGC	TACTGACATT	AAGTGAACGT	CAG	ACG	GCC	GAT	GAA								2349	
					Thr	Ala	Asp	Glu								470	
CAC	ACG	TAT	ATT	CTT	CCA	AAG	AAC	CAA	GTT	GTC	GAG	TTG	CAC	ATC	AAG	2397	
His	Thr	Tyr	Ile	Leu	Pro	Lys	Asn	Gln	Val	Val	Glu	Leu	His	Ile	Lys		
	475				480						485						
GGA	CAG	GCT	TTG	GGA	ATC	GTA	CAC	CCC	CTT	CAT	CTG	CAT	GGC			2439	
Gly	Gln	Ala	Leu	Gly	Ile	Val	His	Pro	Leu	His	Leu	His	Gly				
490					495				500								
GTACGTCTTT	CTCACACTGT	TCCAGCTCCT	ATTCTCTAAC	ACACTCCTGC	GATAC	CAT										2497	
						His											

GCG TTC GAC GTC GTC CAA TTC GGC GAC AAC GCT CCA AAC TAC GTG AAC 2545
Ala Phe Asp Val Val Gln Phe Gly Asp Asn Ala Pro Asn Tyr Val Asn
505 510 515 520

CCT CCG COT AGG GAT GTA GTA GGC GTA ACT GAT GCT GGA GTC CGT ATC 2593
Pro Pro Arg Arg Asp Val Val Gly Val Thr Asp Ala Gly Val Arg Ile
525 530 535

CAG TTC AGA ACC GAT AAC CCG GGC CCT TGG TTC CTC CAT TGC 2635
Gln Phe Arg Thr Asp Asn Pro Gly Pro Trp Phe Leu His Cys
540 545 550

GTATGCTCTT CATCTCCAC CGCTTGTTCT TTACTTATCG TTTACCTTGC GATTTAG 2692

CAC ATT GAT TGG CAC TTG GAA GAA GGA TTT GCT GTAAGTTATT ATTCCTATTC 2745
His Ile Asp Trp His Leu Glu Glu Gly Phe Ala
555 560

CGAAGCATCG GCGAGATGCT AACCAAGGGT GTGTTTAAAG ATG GTA TTC GCC GAA 2800
Met Val Phe Ala Glu
565

GCG CCT GAA GAT ATC AAG AAA GGC TCT CAG AGT GTC AAG CCT GAC GGA 2848
Ala Pro Glu Asp Ile Lys Lys Gly Ser Gln Ser Val Lys Pro Asp Gly
570 575 580

CAA TGG AAG AAA CTA TGC GAG AAG TAT GAG AAG TTG CCT GAA GCA CTG 2896
Gln Trp Lys Lys Leu Cys Glu Lys Tyr Glu Lys Leu Pro Glu Ala Leu
585 590 595

CAG TGAAGTTGCA GTTGTTCCTT ATTCGGGAAC TGGCTCACTA TTCCTTTTGC 2949
Gln

ATAATTCGGA CTTTATTTTT GGGACATTAT TGGACTATGG ACTTGTTCGT CACACCCTCG 3009

CTCACTGTGT CCCTCGTTGA GTACCTATAC TCTATTCGTA TAGTGGGAAT ATGGAATATC 3069

GGATGTAATA AATGCTCGTG CGTTTGGTGC TCGAAATGGG GTAGGACT 3117

(2) INFORMATION FOR SEQ ID NO:4:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 599 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

Met Ala Arg Ser Thr Thr Ser Leu Phe Ala Leu Ser Leu Val Ala Ser
1 5 10 15

Ala Phe Ala Arg Val Val Asp Tyr Gly Phe Asp Val Ala Asn Gly Ala
20 25 30

Val Ala Pro Asp Gly Val Thr Arg Asn Ala Val Leu Val Asn Gly Arg
35 40 45

Phe Pro Gly Pro Leu Ile Thr Ala Asn Lys Gly Asp Thr Leu Lys Ile
50 55 60

Thr Val Arg Asn Lys Leu Ser Asp Pro Thr Met Arg Arg Ser Thr Thr
65 70 75 80

Ile His Trp His Gly Leu Leu Gln His Arg Thr Ala Glu Glu Asp Gly
 85 90 95
 Pro Ala Phe Val Thr Gln Cys Pro Ile Pro Pro Gln Glu Ser Tyr Thr
 100 105 110
 Tyr Thr Met Pro Leu Gly Glu Gln Thr Gly Thr Tyr Trp Tyr His Ser
 115 120 125
 His Leu Ser Ser Gln Tyr Val Asp Gly Leu Arg Gly Pro Ile Val Ile
 130 135 140
 Met Asp Pro His Asp Pro Tyr Arg Asn Tyr Tyr Asp Val Asp Asp Glu
 145 150 155 160
 Arg Thr Val Phe Thr Leu Ala Asp Trp Tyr His Thr Pro Ser Glu Ala
 165 170 175
 Ile Ile Ala Thr His Asp Val Leu Lys Thr Ile Pro Asp Ser Gly Thr
 180 185 190
 Ile Asn Gly Lys Gly Lys Tyr Asp Pro Ala Ser Ala Asn Thr Asn Asn
 195 200 205
 Thr Thr Leu Glu Asn Leu Tyr Thr Leu Lys Val Lys Arg Gly Lys Arg
 210 215 220
 Tyr Arg Leu Arg Ile Ile Asn Ala Ser Ala Ile Ala Ser Phe Arg Phe
 225 230 235 240
 Gly Val Gln Gly His Lys Cys Thr Ile Ile Glu Ala Asp Gly Val Leu
 245 250 255
 Thr Lys Pro Ile Glu Val Asp Ala Phe Asp Ile Leu Ala Gly Gln Arg
 260 265 270
 Tyr Ser Cys Ile Leu Lys Ala Asp Gln Asp Pro Asp Ser Tyr Trp Ile
 275 280 285
 Asn Ala Pro Ile Thr Asn Val Leu Asn Thr Asn Val Gln Ala Leu Leu
 290 295 300
 Val Tyr Glu Asp Asp Lys Arg Pro Thr His Tyr Pro Trp Lys Pro Phe
 305 310 315 320
 Leu Thr Trp Lys Ile Ser Asn Glu Ile Ile Gln Tyr Trp Gln His Lys
 325 330 335
 His Gly Ser His Gly His Lys Gly Lys Gly His His His Lys Val Arg
 340 345 350
 Ala Ile Gly Gly Val Ser Gly Leu Ser Ser Arg Val Lys Ser Arg Ala
 355 360 365
 Ser Asp Leu Ser Lys Lys Ala Val Glu Leu Ala Ala Ala Leu Val Ala
 370 375 380
 Gly Glu Ala Glu Leu Asp Lys Arg Gln Asn Glu Asp Asn Ser Thr Ile
 385 390 395 400
 Val Leu Asp Glu Thr Lys Leu Ile Pro Leu Val Gln Pro Gly Ala Pro
 405 410 415
 Gly Gly Ser Arg Pro Ala Asp Val Val Val Pro Leu Asp Phe Gly Leu
 420 425 430
 Asn Phe Ala Asn Gly Leu Trp Thr Ile Asn Asn Val Ser Tyr Ser Pro

435					440					445					
Pro	Asp	Val	Pro	Thr	Leu	Leu	Lys	Ile	Leu	Thr	Asp	Lys	Asp	Lys	Val
	450					455					460				
Asp	Ala	Ser	Asp	Phe	Thr	Ala	Asp	Glu	His	Thr	Tyr	Ile	Leu	Pro	Lys
465					470					475					480
Asn	Gln	Val	Val	Glu	Leu	His	Ile	Lys	Gly	Gln	Ala	Leu	Gly	Ile	Val
				485					490					495	
His	Pro	Leu	His	Leu	His	Gly	His	Ala	Phe	Asp	Val	Val	Gln	Phe	Gly
			500					505					510		
Asp	Asn	Ala	Pro	Asn	Tyr	Val	Asn	Pro	Pro	Arg	Arg	Asp	Val	Val	Gly
		515					520					525			
Val	Thr	Asp	Ala	Gly	Val	Arg	Ile	Gln	Phe	Arg	Thr	Asp	Asn	Pro	Gly
	530					535					540				
Pro	Trp	Phe	Leu	His	Cys	His	Ile	Asp	Trp	His	Leu	Glu	Glu	Gly	Phe
545					550					555					560
Ala	Met	Val	Phe	Ala	Glu	Ala	Pro	Glu	Asp	Ile	Lys	Lys	Gly	Ser	Gln
				565					570					575	
Ser	Val	Lys	Pro	Asp	Gly	Gln	Trp	Lys	Lys	Leu	Cys	Glu	Lys	Tyr	Glu
			580					585					590		
Lys	Leu	Pro	Glu	Ala	Leu	Gln									
			595												

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

Ala Val Arg Asn Tyr Lys Phe Asp Ile Lys Asn Val Asn Val Ala Pro
1 5 10 15
Asp Gly Phe Gln Arg Pro Ile Val Ser Val
20 25

(2) INFORMATION FOR SEQ ID NO:6:

(1) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 20 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

Ser Gln Tyr Val Asp Gly Leu Arg Gly Pro Leu Val Ile Tyr Asp Pro
1 5 10 15
Asp Asp Asp His
20

(2) INFORMATION FOR SEQ ID NO:7:

(1) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 26 amino acids

(B) TYPE: amino acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

Ser Arg Tyr Asx Val Asx Asx Ala Ser Thr Val Val Met Leu Glu Asx
 1 5 10 15

Trp Tyr Arg Thr Pro Ala Xaa Val Leu Glu
 20 25

(2) INFORMATION FOR SEQ ID NO:8:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 25 amino acids
 (B) TYPE: amino acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

Ser Leu Gly Pro Thr Pro Asn Tyr Val Asn Pro Xaa Ile Arg Asp Val
 1 5 10 15

Val Arg Val Gly Gly Thr Thr Val Val
 20 25

(2) INFORMATION FOR SEQ ID NO:9:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 25 amino acids
 (B) TYPE: amino acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

Gly Leu Ala Leu Val Phe Ala Glu Ala Pro Ser Gln Ile Arg Gln Gly
 1 5 10 15

Val Gln Ser Val Gln Pro Asp Asp Ala
 20 25

(2) INFORMATION FOR SEQ ID NO:10:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 14 amino acids
 (B) TYPE: amino acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

Ile Arg Tyr Val Gly Gly Pro Ala Val Xaa Arg Ser Val Ile
1 5 10

(2) INFORMATION FOR SEQ ID NO:11:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 6 amino acids

(B) TYPE: amino acid

(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

Ile Leu Ala Asn Pro Ala
1 5

(2) INFORMATION FOR SEQ ID NO:12:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 8 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

Tyr Glu Ala Pro Ser Leu Pro Thr
1 5

(2) INFORMATION FOR SEQ ID NO:13:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1912 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(vi) ORIGINAL SOURCE:
(A) ORGANISM: Rhizoctonia laccase

(ix) FEATURE:
(A) NAME/KEY: CDS
(E) LOCATION: 85..1671

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:

CTAACGCTTG GTGCGGAGCT CGGATCCACT AGTAACGGCG GCCAGTGTGC TGGAAATTCGC	60
GGCGCGCTGG ACACCTCCTT CAAG ATG CTT TCT AGC ATT ACC CTC CTA CCT	111
Met Leu Ser Ser Ile Thr Leu Leu Pro	
1 5	
TTG CTC GCT GCG GTC TCA ACC GCG GCC TTT GCT GCC GTC CGC AAC TAT	159
Leu Leu Ala Ala Val Ser Thr Pro Ala Phe Ala Ala Val Arg Asn Tyr	
10 15 20 25	
AAG TTC GAC ATC AAG AAC GTC AAT GTC GCT CCC GAT GGC TTT CAG CGC	207
Lys Phe Asp Ile Lys Asn Val Asn Val Ala Pro Asp Gly Phe Gln Arg	
30 35 40	
TCT ATC CTC TCC GTC AAC GGT TTA GTT CCT GGC ACG TTG ATC ACG GCC	255
Ser Ile Val Ser Val Asn Gly Leu Val Pro Gly Thr Leu Ile Thr Ala	
45 50 55	
AAC AAG GGT GAC ACC TTG CGC ATT AAT GTC ACG AAT CAA CTC ACG GAC	303
Asn Lys Gly Asp Thr Leu Arg Ile Asn Val Thr Asn Gln Leu Thr Asp	
60 65 70	
CCT AGT ATG CGT CGT GCC ACA ACG ATT CAT TGG CAT GGA TTG TTC CAA	351
Pro Ser Met Arg Arg Ala Thr Thr Ile His Trp His Gly Leu Phe Gln	
75 80 85	

GCT ACT ACC GCC GAC GAG GAT GGC CCC GCA TTC GTC ACG CAA TGC CCT Ala Thr Thr Ala Asp Glu Asp Gly Pro Ala Phe Val Thr Gln Cys Pro 90 95 100 105	399
ATT GCG CAA AAT TTG TCC TAT ACA TAC GAG ATC CCA TTG CGC GGC CAA Ile Ala Gln Asn Leu Ser Tyr Thr Tyr Glu Ile Pro Leu Arg Gly Gln 110 115 120	447
ACA GGA ACC ATG TGG TAT CAC GCC CAT CTT GCG AGT CAA TAT GTC GAT Thr Gly Thr Met Trp Tyr His Ala His Leu Ala Ser Gln Tyr Val Asp 125 130 135	495
GGA TTG CGA GGC CCT TTG GTC ATC TAT GAT CCA AAC GAC CCA CAC AAG Gly Leu Arg Gly Pro Leu Val Ile Tyr Asp Pro Asn Asp Pro His Lys 140 145 150	543
TCG CGC TAC GAC GTG GAT GAT GCG AGC ACA GTA GTC ATG CTT GAG GAC Ser Arg Tyr Asp Val Asp Ala Ser Thr Val Met Leu Glu Asp 155 160 165	591
TGG TAC CAT ACT CCG GCA CCC GTT CTA GAA AAG CAA ATG TTC TCG ACT Trp Tyr His Thr Pro Ala Pro Val Leu Glu Lys Gln Met Phe Ser Thr 170 175 180 185	639
AAT AAC ACC GCT CTG CTC TCT CCT GTT CCG GAC TCG GGT CTT ATC AAT Asn Asn Thr Ala Leu Leu Ser Pro Val Pro Asp Ser Gly Leu Ile Asn 190 195 200	687
GGC AAA GGG CGC TAT GTG GGC GGT CCC GCA GTT CCC CGG TCA GTA ATC Gly Lys Gly Arg Tyr Val Gly Gly Pro Ala Val Pro Arg Ser Val Ile 205 210 215	735
AAC GTA AAA CGT GGG AAA CGA TAT CGC TTG CGC GTA ATC AAC GCT TCT Asn Val Lys Arg Gly Lys Arg Tyr Arg Leu Arg Val Ile Asn Ala Ser 220 225 230	783
GCT ATC GGG TCG TTT ACC TTT TCG ATC GAA GGA CAT AGT CTG ACT GTC Ala Ile Gly Ser Phe Thr Phe Ser Ile Glu Gly His Ser Leu Thr Val 235 240 245	831
ATT GAG GCC GAT GGG ATC CTG CAC CAG CCC TTG GCT GTT GAC AGC TTC Ile Glu Ala Asp Gly Ile Leu His Gln Pro Leu Ala Val Asp Ser Phe 250 255 260 265	879
CAG ATT TAC GCT GGA CAA CGC TAC TCT GTC ATC GTT GAA GCC AAC CAA Gln Ile Tyr Ala Gly Gln Arg Tyr Ser Val Ile Val Glu Ala Asn Gln 270 275 280	927
ACC GCC GCC AAC TAC TGG ATT CGT GCA CCA ATG ACC GTT GCA GGA GCC Thr Ala Ala Asn Tyr Trp Ile Arg Ala Pro Met Thr Val Ala Gly Ala 285 290 295	975
GGA ACC AAT GCA AAC TTG GAC CCC ACC AAT GTC TTT GCC GTA TTG CAC Gly Thr Asn Ala Asn Leu Asp Pro Thr Asn Val Phe Ala Val Leu His 300 305 310	1023
TAC GAG GGA GCG CCC AAC GCC GAA CCC ACG ACG GAA CAA GGC AGT GCT Tyr Glu Gly Ala Pro Asn Ala Glu Pro Thr Thr Gln Gly Ser Ala 315 320 325	1071
ATC GGT ACT GCA CTC GTT GAA GAG AAC CTC CAT GCG CTC ATC AAC CCT Ile Gly Thr Ala Leu Val Glu Glu Asn Leu His Ala Leu Ile Asn Pro 330 335 340 345	1119
GGC GCT CCG GGC GGC TCC GCT CCC GCA GAC GTT TCC CTC AAT CTT GCA Gly Ala Pro Gly Gly Ser Ala Pro Ala Asp Val Ser Leu Asn Leu Ala 350 355 360	1157

ATT GGG CGC AGC ACA GTT GAT GGG ATT CTT AGG TTC ACA TTT AAT AAC	1215
Ile Gly Arg Ser Thr Val Asp Gly Ile Leu Arg Phe Thr Phe Asn Asn	
365 370 375	
ATC AAG TAC GAG GCT COT TCG TTG CCC ACG CTC TTG AAG ATT TTG GCA	1263
Ile Lys Tyr Glu Ala Pro Ser Leu Pro Thr Leu Leu Lys Ile Leu Ala	
380 385 390	
AAC AAT GCG AGC AAT GAC GCC GAT TTC ACG CCA AAT GAG CAC ACT ATC	1311
Asn Asn Ala Ser Asn Asp Ala Asp Phe Thr Pro Asn Glu His Thr Ile	
395 400 405	
GTA TTG CCA CAC AAT AAA GTT ATC GAG CTC AAT ATC ACC GGA GGT GCA	1359
Val Leu Pro His Asn Lys Val Ile Glu Leu Asn Ile Thr Gly Gly Ala	
410 415 420 425	
GAC CAC CCT ATC CAT CTC CAC GGC CAT GTG TTT GAT ATC GTC AAA TCA	1407
Asp His Pro Ile His Leu His Gly His Val Phe Asp Ile Val Lys Ser	
430 435 440	
CTC GGT GGT ACC CCG AAC TAT GTC AAC CCG CCA CCG AGG GAC GTA GTT	1455
Leu Gly Gly Thr Pro Asn Tyr Val Asn Pro Pro Arg Arg Asp Val Val	
445 450 455	
CGT GTC GGA GGC ACC GGT GTG GTA CTC CGA TTC AAG ACC GAT AAC CCA	1503
Arg Val Gly Gly Thr Gly Val Val Leu Arg Phe Lys Thr Asp Asn Pro	
460 465 470	
GGC CCA TGG TTT GTT CAC TGC CAC ATT GAC TGG CAC TTG GAG GCT GGG	1551
Gly Pro Trp Phe Val His Cys His Ile Asp Trp His Leu Glu Ala Gly	
475 480 485	
CTC GCA CTT GTC TTT GCC GAG GCC CCC AGC CAG ATT CCG CAC GGT GTC	1599
Leu Ala Leu Val Phe Ala Glu Ala Pro Ser Gln Ile Arg Gln Gly Val	
490 495 500 505	
CAG TCG GTC CAG CCC AAC AAT GCC TGG AAC CAG CTC TGC CCC AAG TAC	1647
Gln Ser Val Gln Pro Asn Asn Ala Trp Asn Gln Leu Cys Pro Lys Tyr	
510 515 520	
GCG GCT CTT CCT CCC GAT TFG CAG T	1672
Ala Ala Leu Pro Pro Asp Leu Gln	
525	

(2) INFORMATION FOR SEQ ID NO:14:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 529 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:14:

Met Leu Ser Ser Ile Thr Leu Leu Pro Leu Leu Ala Ala Val Ser Thr
 1 5 10 15
 Pro Ala Phe Ala Ala Val Arg Asn Tyr Lys Phe Asp Ile Lys Asn Val
 20 25 30
 Asn Val Ala Pro Asp Gly Phe Gln Arg Ser Ile Val Ser Val Asn Gly
 35 40 45
 Leu Val Pro Gly Thr Leu Ile Thr Ala Asn Lys Gly Asp Thr Leu Arg
 50 55 60

Ile Asn Val Thr Asn Gln Leu Thr Asp Pro Ser Met Arg Arg Ala Thr
 65 70 75 80
 Thr Ile His Trp His Gly Leu Phe Gln Ala Thr Thr Ala Asp Glu Asp
 85 90 95
 Gly Pro Ala Phe Val Thr Gln Cys Pro Ile Ala Gln Asn Leu Ser Tyr
 100 105 110
 Thr Tyr Glu Ile Pro Leu Arg Gly Gln Thr Gly Thr Met Trp Tyr His
 115 120 125
 Ala His Leu Ala Ser Gln Tyr Val Asp Gly Leu Arg Gly Pro Leu Val
 130 135 140
 Ile Tyr Asp Pro Asn Asp Pro His Lys Ser Arg Tyr Asp Val Asp Asp
 145 150 155 160
 Ala Ser Thr Val Val Met Leu Glu Asp Trp Tyr His Thr Pro Ala Pro
 165 170 175
 Val Leu Glu Lys Gln Met Phe Ser Thr Asn Asn Thr Ala Leu Leu Ser
 180 185 190
 Pro Val Pro Asp Ser Gly Leu Ile Asn Gly Lys Gly Arg Tyr Val Gly
 195 200 205
 Gly Pro Ala Val Pro Arg Ser Val Ile Asn Val Lys Arg Gly Lys Arg
 210 215 220
 Tyr Arg Leu Arg Val Ile Asn Ala Ser Ala Ile Gly Ser Phe Thr Phe
 225 230 235 240
 Ser Ile Glu Gly His Ser Leu Thr Val Ile Glu Ala Asp Gly Ile Leu
 245 250 255
 His Gln Pro Leu Ala Val Asp Ser Phe Gln Ile Tyr Ala Gly Gln Arg
 260 265 270
 Tyr Ser Val Ile Val Glu Ala Asn Gln Thr Ala Ala Asn Tyr Trp Ile
 275 280 285
 Arg Ala Pro Met Thr Val Ala Gly Ala Gly Thr Asn Ala Asn Leu Asp
 290 295 300
 Pro Thr Asn Val Phe Ala Val Leu His Tyr Glu Gly Ala Pro Asn Ala
 305 310 315 320
 Glu Pro Thr Thr Glu Gln Gly Ser Ala Ile Gly Thr Ala Leu Val Glu
 325 330 335
 Glu Asn Leu His Ala Leu Ile Asn Pro Gly Ala Pro Gly Gly Ser Ala
 340 345 350
 Pro Ala Asp Val Ser Leu Asn Leu Ala Ile Gly Arg Ser Thr Val Asp
 355 360 365
 Gly Ile Leu Arg Phe Thr Phe Asn Asn Ile Lys Tyr Glu Ala Pro Ser
 370 375 380
 Leu Pro Thr Leu Leu Lys Ile Leu Ala Asn Asn Ala Ser Asn Asp Ala
 385 390 395 400
 Asp Phe Thr Pro Asn Glu His Thr Ile Val Leu Pro His Asn Lys Val
 405 410 415
 Ile Glu Leu Asn Ile Thr Gly Gly Ala Asp His Pro Ile His Leu His

	420		425		430														
Gly	His	Val	Phe	Asp	Ile	Val	Lys	Ser	Leu	Gly	Gly	Thr	Pro	Asn	Tyr				
	435						440					445							
Val	Asn	Pro	Pro	Arg	Arg	Asp	Val	Val	Arg	Val	Gly	Gly	Thr	Gly	Val				
	450					455					460								
Val	Leu	Arg	Phe	Lys	Thr	Asp	Asn	Pro	Gly	Pro	Trp	Phe	Val	His	Cys				
465					470					475					480				
His	Ile	Asp	Trp	His	Leu	Glu	Ala	Gly	Leu	Ala	Leu	Val	Phe	Ala	Glu				
				485					490						495				
Ala	Pro	Ser	Gln	Ile	Arg	Gln	Gly	Val	Gln	Ser	Val	Gln	Pro	Asn	Asn				
			500					505					510						
Ala	Trp	Asn	Gln	Leu	Cys	Pro	Lys	Tyr	Ala	Ala	Leu	Pro	Pro	Asp	Leu				
	515						520					525							

Gln

What we claim is:

1. A nucleic acid fragment containing a nucleic acid sequence encoding a *Rhizoctonia* laccase which functions optimally at
5 pH between about 6.0 and 8.5.
2. The fragment of Claim 1 which comprises a sequence encoding a *Rhizoctonia solani* laccase.
- 10 3. The fragment of Claim 1 which comprises a nucleic acid sequence encoding the amino acid sequence depicted in SEQ ID NO. 2.
4. The fragment of Claim 1 which comprises a nucleic acid
15 sequence encoding the amino acid sequence depicted in SEQ ID NO. 4.
5. The fragment of Claim 1, which comprises a nucleic acid sequence encoding a protein containing one or more of the
20 amino acid sequences depicted in SEQ. ID NOS. 5, 6, 7, 8, 9, 10, 11, or 12.
6. The fragment of Claim 1 which comprises a nucleic acid sequence encoding the amino acid sequence depicted in SEQ ID
25 NO. 14.
7. The fragment of Claim 1, which comprises the nucleic acid sequence depicted in SEQ ID NO. 1.
- 30 8. The fragment of Claim 1, which comprises the nucleic acid sequence depicted in SEQ. ID. NO. 3.

9. The fragment of Claim 1, which comprises the nucleic acid sequence depicted in SEQ. ID. NO. 13.
10. The fragment of Claim 1, which comprises the nucleic acid sequence contained in NRRL B-21141.
11. The fragment of Claim 1, which comprises the nucleic acid sequence contained in NRRL B-21142.
12. The fragment of Claim 1, which comprises the nucleic acid sequence encoding the laccase produced by RS 22.
13. The fragment of Claim 1, which comprises the nucleic acid sequence contained in NRRL B-21156.
14. A substantially pure *Rhizoctonia* laccase enzyme which functions optimally at a pH between about 6.0-8.5.
15. The enzyme of Claim 14 which is a *Rhizoctonia solani* laccase.
16. The enzyme of Claim 14 which comprises the sequence depicted in SEQ ID NO. 2, or a sequence with at least 80% homology thereto.
17. The enzyme of Claim 14 which comprises the sequence depicted in SEQ ID NO 4, or a sequence with at least 80% homology thereto.
18. The enzyme of Claim 14 which comprises one or more of the peptide sequences depicted in SEQ ID NOS.5, 6, 7,

8, 9, 10, 11 or 12, or a sequence with at least 80% homology to one or more of these peptides.

19. The enzyme of Claim 14 which comprises the sequence
5 depicted in SEQ ID NO 14, or a sequence with at least 80% homology thereto.

20. A recombinant vector comprising a nucleic acid fragment containing a nucleic acid sequence encoding a *Rhizoctonia*
10 laccase which functions optimally at pH between about 6.0-8.5.

21. The vector of Claim 20 in which the fragment is operably linked to a promoter sequence.

15

22. The vector of Claim 21 in which the promoter is a fungal or yeast promoter.

23. The vector of Claim 22 in which the promoter is the
20 TAKA amylase promoter of *Aspergillus oryzae*.

24. The vector of Claim 22 in which the promoter is the glucoamylase (gluA) promoter of *Aspergillus niger* or *Aspergillus awamsii*.

25

25. The vector of Claim 21 which also comprises a selectable marker.

26. The vector of Claim 25 in which the selectable marker
30 is the amdS marker of *Aspergillus nidulans* or *Aspergillus oryzae*.

27. The vector of Claim 25 in which the selectable marker is the pyrG marker of *Aspergillus nidulans*, *Aspergillus niger*, *Aspergillus awamorii*, or *Aspergillus oryzae*.
- 5 28. The vector of Claim 21 which comprises both the TAKA amylase promoter of *Aspergillus oryzae* and the amdS or pyrG marker of *Aspergillus nidulans* or *Aspergillus oryzae*.
29. A host cell comprising a heterologous nucleic acid
10 fragment containing a nucleic acid sequence encoding a *Rhizoctonia laccase* which functions optimally at pH between about 6.0-8.5.
30. The host cell of Claim 28 which is a fungal cell.
15
31. The host cell of Claim 30 which is an *Aspergillus* cell.
32. The host cell of Claim 29 in which the fragment is integrated into the host cell genome.
20
33. The host cell of Claim 29 in which the fragment is contained on a vector.
34. The host cell of Claim 29 which comprises a fragment
25 containing a sequence encoding the amino acid sequence depicted in SEQ ID NO. 2.
35. The host cell of Claim 29 which comprises a fragment
containing a sequence encoding the amino acid sequence
30 depicted in SEQ ID NO: 4.

36. The host cell of Claim 29 which comprises a fragment containing a sequence encoding the amino acid sequence depicted in SEQ ID NO: 14.
- 5 37. The host cell of Claim 29 which comprises a fragment containing a sequence encoding one or more of the amino acid sequences depicted in SEQ ID NOS.: 5, 6, 7, 8, 9, 10, 11, or 12.
- 10 38. A method for obtaining a laccase enzyme which functions optimally at a pH between about 6.0-8.5 which comprises culturing a host cell comprising a nucleic acid fragment containing a nucleic acid sequence encoding a *Rhizoctonia* laccase enzyme which functions optimally at a pH between
15 about 6.0-8.5, under conditions conducive to expression of the enzyme, and recovering the enzyme from the culture.
39. A method for polymerizing a lignin or lignosulfate substrate in solution which comprises contacting the
20 substrate with a *Rhizoctonia* laccase which functions optimally at a pH between about 6.0-8.5.
40. A method for in situ depolymerization in Kraft pulp which comprises contacting the pulp with a *Rhizoctonia*
25 laccase which functions optimally at a pH between about 6.0-8.5.
41. A method for oxidizing dyes which comprises contacting the dye with a *Rhizoctonia* laccase which functions optimally
30 at a pH between about 6.0-8.5.

42. A method of polymerizing a phenolic compounds which comprises contacting the phenolic compound with a *Rhizoctonia* laccase which functions optimally at a pH between about 6.0-8.5.

5

1 AGCGTCACACCAGACATCGGATGAAACGGAAAGTGATGCGCCATTGACGTCTGCGGC 60
 61 AACCACTGTTCAATCTCGCGAGCTAACATGGCGACGTATAAGAAAGCGGAGAAATGGGC 120
 121 AGATTTCGATATCCCCCTCTCGTCTCGGTTTGGTCTCGGCTTGCCCTCTAATGGCGCGCAC 180
 M A R T 4
 181 CACTTTCCTTGTCTCGGTTTCGGCTCTTTGTTCGGCTGTCTTGGCGCACCGTCGAGTA 240
 4 T F L V S V S L F V S A V L A R T V E Y 24
 241 CGGCTTGAAGATTAGTAGTGGGAGATAGCTCCTGACGGTGTAAAGCGTAATGCGACTTT 300
 24 G L K I S D G E I A P D G V K R N A T L 44
 301 GGgtacgcactccttgtaatccaacaattccaaggttcttgatgcttggtcagTAAATGGA 360
 44 V N G 47
 361 GGGTATCCCGGTCCACTCATTTTTCGCCAACAAGGGGATACTCTCAAAGTCAAGGTCCAA 420
 47 G Y P G P L I F A N K G D T L K V K V Q 67
 421 AACAAGCTCACGAATCCTGAGATGTATCCGACCACTTCCATCGtatgttcggtcgatc 480
 67 N K L T N P E M Y R T S I 81
 481 tactaatcacatccgctcgctaaatatcttgtagCATTTGGCACGGTCTCTTACAACATAGAA 540
 81 H W H G L L Q H R 90

FIG. 1A

541	ACGCCGACGACGACGTCCTTCGTTTCGTCACCTCAGgtaggattctggaagggtggcctga	600
90	N A D D G P S F V T Q	102
601	actctctgttaaccgacaaccccgatgtcaccagTCCCCGATTGTTCCACGCGAGTCGTAT	660
102	C P I V P R E S Y	111
661	ACTTACACCATACCTTCGGACGATCAAAACCGGAACCTATTGGTACCATAGCCACTTGAGT	720
111	T Y T I P L D D Q T G T Y W Y H S H L S	131
721	TCGCAATACGTTGATGGTCTTCGAGGCCCGCTGGTAATCTgtgagtatctcttgacttgtct	780
131	S Q Y V D G L R G P L V I	144
781	actgaaggcaacgagactaaacaagcgctcgattcacagATGgttcgtctccccctttatt	840
144	Y	145
841	tagctctggatcttcoatttctcacgtaatacatgatagATCCCAAGGATCCTCACAGGCG	900
144	D P K D P H R R	152
901	TTTGTATGATGTTGACGATGAGAAAGACCGTCTCTGATCATCGGTGACTGGTATCATGAATC	960
152	L Y D V D D E K T V L I I G D W Y H E S	172
961	GTCCAAGGCAATCCTTGCTTCTGGTAACATTACCCGACAGtaagtgatatacatgcccggtcc	1020
172	S K A I L A S G N I T R Q	185

FIG. 1B

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1021 cagaaaaattctctaaattcatttttaattacagcgcgacccggctcttgcaccatcaacgg 1080
185      R P V S A T I N G
1081 CAAAGGTCGATTGACCCCTGACAAACACTCCTGCCAACCCAGATACCTCTGTACACCCCTCAA 1140
194 K G R F D P D N T P A N P D T L Y T L K
1141 GGTCAGCGAGGGAAGCGCTATCGTCTGCCGTGTCATCAATAGCTCGGAGATCGCTTCGTT 1200
214 V K R G K R Y R L R V I N S S E I A S F
1201 CCGATTCAAGTGTGAAGTCAACAAGGTGACTGTGATTGCTGCCGATGCCGTCTCTACCAA 1260
234 R F S V E G H K V T V I A A D G V S T K
1261 ACCGTATCAGGTCGATGCGTTTGATATCTAGCAGGACAGCGCATAGATTGCGTCgtaag 1320
254 P Y Q V D A F D I L A G Q R I D C V
1321 tgtcgtccgaacccacatctgagctcaagtgtgatacatgcgcgcttatagGTGGAGGC 1380
272      V E A
1381 GAACCAAGAACCCGACACATACCTGGAACAACGCCACCGCTGACCAACGTGCCCAACAAGAC 1440
275 N Q E P D T Y W I N A P L T N V P N K T
1441 CGCTCAGGCTCTCCTCGTTTATGAGGAGGATCGTCCGCCGTACCAACCTCCAAAGGCCCC 1500
295 A Q A L L V Y E E D R R P Y H P P K G P
1501 GTATCGCAAGTGGAGCGTCTCTGAGGCGATCATCAAGTACTGGAATCACAAGCACAAGCA 1560
315 Y R K W S V S E A I I K Y W N H K H K H

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FIG. 1C

3/21

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1561 CGGACGTGGTTGCTGTCTGGACATGGAGGCTCTCAAGGCTCGGATCATCGAGGTAGCCA 1620
335   G R G L L S G H G G L K A R M I E G S H 340

1621 TCATCTGCATTCGCCGACGGTCGTTAAGCGCCAGAATGAGACCACCACTGTTGTATGGA 1680
340   H L H S R S V V K R Q N E T T V V M D 350

1681 CGAGAGCAAGCTCGTTgtaagtaaccataatttaaaagttgggttggttcgaatacttatt 1740
350   E S K L V

1741 tcaacttttcttagCCACTGGAATACCCCGCGCTGCATGCGGGTCTAAACCTGCTGACC 1800
350   P L E Y P G A A C G S K P A D 365

1801 TCGTCTTGGATCTCACTTTTGGTTTGGtatgtagccaaatcgcccatatacaggatactg 1860
365   L V L D L T F G L 374

1861 aatatgttbtgtgcgtgtagAACTTGTCTACCGGGCACTGGATGATCAACGGTATCCCAT 1920
374   N F A T G H W M I N G I P 387

1921 ACGAGTCTCCCAAATCCCCACATTTGCTCAAGATCCTCACTGATGAGACGGGTTACCG 1980
387   Y E S P K I P T L L K I L T D E D G V T 407

1981 AGTCTGACTTgtatgttcccttttcgggtatcttcgtatgcgtgcactgactcgtgctgggt 2040
407   E S D F 411

2041 ggggaatttagCACCAGGAGGAGCACACAGTCATACTCCCGAAGAACAAATGCATCGAAT 2100
411   T K E E H T V I L P K N K C I E 427

```

FIG. 1D

2101 TCAACATCAAGGGAACCTCGGTATTCCTCCATTACGACCCCGTACATCTTCACCGTgtaa 2160
 427 F N I K G N S G I P I T H P V H L H G 446
 2161 gtgcataatcggtggtttacgataactaaggctcatcaactttttagCACACTTGGGATGT 2220
 446 H T W D V 451
 2221 CGTACAATTGGCAACACACCCCAATTAATGTCATCTCCCGTAGGACGTGGTGG 2280
 451 V Q F G N N P P N Y V N P P R R D V V G 471
 2281 CTCACAGATCGGGGTGTGAGGATTCAGTTCAGACCCGACAAATCCAGGACCGTGTCT 2340
 471 S T D A G V R I Q F K T D N P G P W F L 491
 2341 GCACTGgtgcgtcggtcccatcggtccgttatgggttttctaatacgtcccattctattt 2400
 491 H C 493
 2401 tagCCATATTGACTGGCATCTTGAGGAGGTTTCGCAAgtagtactgagaccctaagtgc 2460
 493 H I D W H L E E G F A 504
 2461 tactcgggtcattactgattaccgcgatgtatgcgtctagTGGTGTTCGTAAGCGCCCG 2520
 504 M V F A E A P 511
 2521 AAGCCGTCAAGGAGGTCCAAAGAGCGTGGCCGTGACTCTCAGTGGGAAGGCTGTGTG 2580
 511 E A V K G G P K S V A V D S Q W E G L C 531
 2581 GCAAGTACGACAACTGGCTAAATCAAAATCCGGCCAGCTGTAGGCGTATCGCAGCCACA 2640
 531 G K Y D N W L K S N P G Q L * 545

FIG. 1E

2641 TTGGTGAATGAAAGTTGCATCTTGTTCCTATAACCGGCTCTTATATACGGGTGTCIC 2700
2701 CCAGTAAAGTCGTAGCCCCAATTTCAGCCGAGACAGATATTTAGTGGACTCTTACTCTTGT 2760
2761 GTCCCATTTGACGCACATCGTTGCCATCAACCCTGCTTTTATCGTCCCTCTTTGTAAATTG 2820
2821 TGTTCCTGTAATGTATCG 2838

FIG. 1F

```

1  AAGCTTCGGCATGGATGTCATTTTTGTAATGT 180

181  AAACAAGTTACGAGAAAAACAATAGATCAGTTTTCGCCGAATCGGATGGCTTTGAAACCGGA 240

241  AGTACCGATGGCCGATCCGAGTCGAATGAATTACGCATCTGAAACGGGACCCCTGAGTCG 300

301  AGGCACCCGGCCCTTGGCCGTTAAGTCACCTTGTCGCCCACTAGCACATTTTTCATTCC 360

361  CCCTTTCTTCTCCTCGTCTTCTTCTTCTCTATGGCTCGGTCGACTACTTCACTCTTTG 420
      1      M A R S T S L F 10

421  CACTGTCTCTGGCCGCACCGCCCTTGGCTCGAGTCGTTGACTATGGGTTTGATGTGGCTA 480
      10 A L S L A A P A L A R V V D Y G F D V A 30

481  ATGGGGCCAGTTGCTCCGGATGGTGTAAACAAGGAACGGGTTCTCGgtgagbttagctgtaa 540
      30 N G A V A P D G V T R N A V L 45

541  gatggtgtatatgctggttgcctaacgggaatgtcagTCAATGGTCGCTTCCCCTGTCCA 600
      45      V N G R F P G P 53

601  TTGATCACCGCCACAAGGGGATACACTTAAATCACCGTCCGGAATAAATCTCCGAT 660
      53 L I T A N K G D T L K I T V R N K L S D 73

```

FIG. 2A

```

661 CCAACTATGCGAAGGAGCAGACCATCgttagtacttccccctcatctgtcttgaacttt 720
73 P T M R R S T T I 82

721 ctcatcttttttgaagCACTGGCACCgGTCTGCTCCAAACAGGACGGCAGAGAAGATGG 780
82 H W H G L L Q H R T A E D G 97

781 CCCGGCCTTGTAAACCCAGgtatgccttatccctatcgctgctctgtccccgcgtccttcc 840
97 P A F V T Q 103

841 ctgactcgggcgattctagTGCCCGATTCTCCGCAAGAAATCGTACACCTATACGATGCC 900
103 C P I P P Q E S Y T Y T M P 117

901 GCTCGGCGAACAGACCGGACGTATTGGTACCACAGCCCACTTGAGCTCCCAAGTATGTGGA 960
117 L G E Q T G T Y W Y H S H L S S Q Y V D 137

961 CGGGTTGCGTGGCCCATCGTTATTgtaagtcttcatttaacctattcttggctatgg 1020
137 G L R G P I V I 145

1021 ctgattgtgacgtcgtggtagATGgttcgtggcttccacaagaagtcagcagcccttga 1080
145 Y 145

1081 agctaaactttattccagACCCCCACGACCCGTACAGAAACTACTATGATGTCGACGACGA 1140
145 D P H D P Y R N Y Y D V D D E 160

1141 GCGTACCGTCTTTTACCTTAGCAGACTGGTACCACACGCCGTGCGAGGCTATCATTCGCCAC 1200
160 R T V F T L A D W Y H T P S E A I I A T 180

```

FIG. 2B


```

1201 CCACGATGTCCTGAAACgtacgcgttaatccttctagctttcttctccttgggtcacttt 1260
180   H D V L K T
1261 ctatcagGATCCCCGACTCGGGTACGATCAACGGCAAGCAAAATACGATCCTGCTTCGG 1320
185   I P D S G T I N G K G K Y D P A S
1321 CTAACACCAACAACACGACACTCGAGAACCTCTACACTCTCAAGTCAAAACGGCGCAAGC 1380
202 A N T N T T L E N L Y T L K V K R G K
1381 GGTATCGCCTGAGGATTATCAACGCCCTCGGCCATCGCTTCGTCGGTTCGGCGTGCAGG 1440
222 R Y R L R I I N A S A I A S F R F G V Q
1441 GCCACAAGTGCACGATCATCGAGGCTGATGGCGTCCCTCACCAACCGATCGAGTCGATG 1500
242 G H K C T I I E A D G V L T K P I E V D
1501 CGTTTGATAATTCTAGCAGGCCAGAGGTATAGCTGCATCGtaagtctacctatgccttggt 1560
262 A F D I L A G Q R Y S C I
1561 gtggagataagaacctgactgaatgtatgcgctccaatagTTGAAGGCCGACCAAGATCC 1620
275   L K A D Q D P
1621 TGATTCCTACTGGATAAATGCGCCAATCACAACGTTCTCAACACCAACGTCAGGCATT 1680
282   D S Y W I N A P I T N V L N T N V Q A L
1681 GCTAGTGTATGAAGATGACAAGCGTCTCTACTACTACCCCTGGAAGCCGTTTGTGACATG 1740
302   L V Y E D D K R P T H Y P W K P F L T W

```

FIG. 2C

```

1741 GAAGATATCAAAATGAAATCATTCAGTACTGGCAGCACAAAGCAGGTCGACGGTCACAA 1800
322 K I S N E I I Q Y W Q H K H G S H G H K 342
1801 GGGAAAGGGCATCATATAAAGTCCGGCCCATTTGGAGGTGTATCCGGTTGAGCTCCAG 1860
342 G K G H H K V R A I G G V S G L S S R 362
1861 GGTAAAGAGCCGGGAGTGAACCTATTCGAAGAAGGCTGTTCGAGTTGGCTGCTGCACTCGT 1920
349 V K S R A S D L S K K A V E L A A A L V 349
1921 TCCGGGTGAGCCGAGTTGGACAAGAGGAGGAGGATAATTCGACTATTTGTATTGGA 1980
349 A G E A E L D K R Q N E D N S T I V L D 361
1981 TGAGACCAAGCTTATTgttaagtcaccttaatttttttcggtgtcaoggaagctaaccgcg 2040
361 E T K L I 361
2041 taatagCCGTTGTTCAACCTGGTGCACCGGGCGGCTCCAGACCAGCTGACGTGCTGTC 2100
361 P L V Q P G A P G G S R P A D V V V 379
2101 CCTCTGGACTTTGGCCCTCgtatgtggcttcttggtatttcgtccggaatgcaaaactgattt 2160
379 P L D F G L 385
2161 ggggtgggctatagaACTTTGCCAACGGACTGTGGACGATAAACAATGTCCTCTACTCCCC 2220
385 N F A N G L W T I N N V S Y S P 401
2221 TCCGGATGTCCCTACTCTCCTCAAGATCTTGACCGACAAAGACAAAGTCGACGCTTCTGA 2280
401 P D V P T L L K I L T D K D K V D A S D 421

```

FIG. 2D

```

2281 CTTgtaggttccctcttcttcttcttcaactagctactgacattaaagtgaacgtcagCACG 2340
421 F T
2341 GCCGATGAACACACGTATATTCCTTCCAAAGAACCAAGTTGTCGAGTTGCACATCAAGGGA 2400
423 A D E H T Y I L P K N Q V V E L H I K G 453
2401 CAGGCTTTGGGAATCGTACACCCCTTCATCTGTCATGCGCgtacgtcttctcacactggt 2460
453 Q A L G I V H P L H L H G 466
2461 ccagctcctattctctaacacacactcctgcgatagCATGCGTTGACGTCGTCCAAATTCGG 2520
466 H A F D V V Q F G 475
2521 CGACAAACGCTCCAAACTACGTGAACCCCTCCGCTAGGATGTAGTGGCGTAACCTGATGC 2580
475 D N A P N Y V N P P R R D V V G V T D A 495
2581 TGGAGTCCGTATCCAGTTCAGAACCGATACCCGGCCCTTGCTTCCTCCATTGgtatgc 2640
495 G V R I Q F R T D N P G P W F L H C 513
2641 tcttcacatctcccacgcttggttcttacttattggtttaccttgcgatttagCCACATTGA 2700
513 H I D 516
2701 TTGGCACTTGGAAGAAGGATTTCGTAGtaagttatttccctattccgaagcatcg99ga 2760
516 W H L E E G F A 524
2761 gatgctaaccaagggtgtgtttaaagTGGTATTCGCCGAAGCGCCTGAAGATATCAAGAA 2820
524 M V F A E A P E D I K K 536

```

FIG. 2E

```

2821 AGGCTCTCAGAGTGTCAAGCCCTGACGGACAATGGAAGAACTATGCGAGAAGTATGAGAA 2880
536  G S Q S V K P D G Q W K K L C E K Y E K 556

2881 GTTCCCTGAAGCACTGCAGTGAAGTTGCAGTGTTCCTCCATTCGGAACTGGCTCACTAT 2940
556  L P E A L Q * 562

2941 TCCTTTTGCATAATTTCGGACTTTTATTTTGGGACATTATTGGACTATGCAATTGTGTGTC 3000

3001 ACACCGCGGAACATAAGCCGAATTC

```

FIG. 2F

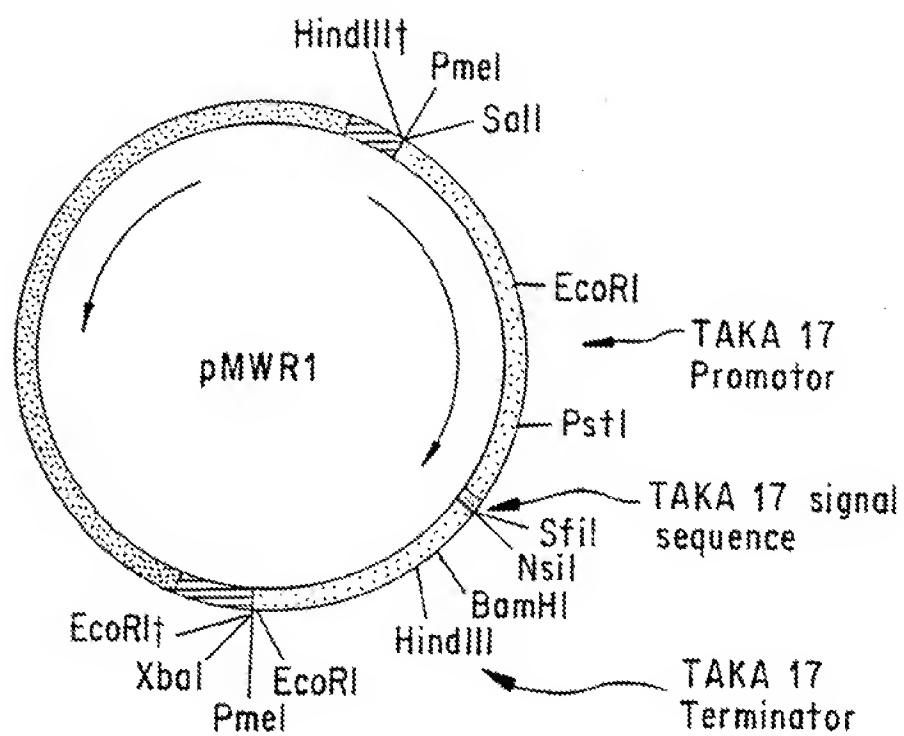


FIG. 3

5'	87	96	105	114	123	132
ATG CTT TCT AGC ATT ACC CTC CTA CCT TTG CTC GCT GCG GTC TCA ACC CCC GCC						

M L S S I T L L P L L A A V S T P A						

141	150	159	168	177	186	
TTF GCT GCC GTC CGC AAC TAT AAG TTC GAC ATC AAG AAC GTC AAT GTC GCT CCC						

F A A V R N Y K F D I K N V A P						

195	204	213	222	231	240	
GAT GGC TTT CAG CGC TCT ATC GTC TCC GTC AAC GGT TTA GTT CCT GGC ACG TTG						

D G F Q R S I V S V N G L V P G T L						

249	258	267	276	285	294	
ATC ACG GCC AAC AAG GGT GAC ACC TTG CGC ATT AAT GTC ACG AAT CAA CTC ACG						

I T A N K G D T L R I N V T N Q L T						

303	312	321	330	339	348	
GAC CCT AGT ATG CGT CGT GCC ACA ACG ATT CAT TGG CAT GGA TTG TTC CAA GCT						

D P S M R R A T T I H W H G L F Q A						

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FIG. 4A

357	366	375	384	393	402
ACT ACC GCC GAC GAG GAT GGC CCC GCA TTC GTC ACG CAA TGC CCT ATT GCG CAA					
---	---	---	---	---	---
T T A D E D G P A F V T Q C P I A Q					
411	420	429	438	447	456
AAT TTG TCC TAT ACA TAC GAG ATC CCA TTG CGC GGC CAA ACA GGA ACC ATG TGG					
---	---	---	---	---	---
N L S Y T Y E I P L R G Q T G T M W					
465	474	483	492	501	510
TAT CAC GCC CAT CTT GCG AGT CAA TAT GTC GAT GGA TTG CGA GGC CCT TTG GTC					
---	---	---	---	---	---
Y H A H L A S Q Y V D G L R G P L V					
519	528	537	546	555	564
ATC TAT GAT CCA AAC GAC CCA CAC AAG TCG CGC TAC GAC GTG GAT GAT GCG AGC					
---	---	---	---	---	---
I Y D P N D P H K S R Y D V D D A S					
573	582	591	600	609	618
ACA GTA GTC ATG CTT GAG GAC TGG TAC CAT ACT CCG GCA CCC GTT CTA GAA AAG					
---	---	---	---	---	---
T V V M L E D W Y H T P A P V L E K					

FIG. 4B

CAA	ATG	TTC	TCG	ACT	AAT	AAC	ACC	GCT	CTG	CTC	TCT	TCT	CCT	GTT	CCG	GAC	TCG	GGT	672
---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---
Q	N	F	S	T	N	N	T	A	L	L	S	P	P	V	P	D	S	G	
CTT	ATC	AAT	GGC	AAA	GGG	CGC	TAT	GTG	GGC	GGT	CCC	GCA	GTT	CCC	CGG	TCA	GTA	726	
---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---
L	I	N	G	K	G	R	Y	V	G	G	P	A	V	P	R	S	V		
ATC	AAC	GTA	AAA	CGT	GGG	AAA	CGA	TAT	CGC	TTG	CGC	GTA	ATC	AAC	GCT	TCT	GCT	780	
---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---
I	N	V	K	R	G	K	R	Y	R	L	R	V	I	N	A	S	A		
ATC	GGG	TCG	TTT	ACC	TTT	TCG	ATC	GAA	GGA	CAT	AGT	CTG	ACT	GTC	ATT	GAG	GCC	834	
---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---
I	G	S	F	T	F	S	I	E	G	H	S	L	T	V	I	E	A		
GAT	GGG	ATC	CTG	CAC	CAG	CCC	TTG	GCT	GTT	GAC	AGC	TTC	CAG	ATT	TAC	GCT	GGA	888	
---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---
D	G	I	L	H	Q	P	L	A	V	D	S	F	Q	I	Y	A	G		

FIG. 4C

897	906	915	924	933	942
CAA CGC TAC TCT GTC ATC GTT GAA GCC AAC CAA ACC GCC AAC TAC TGG ATT					
Q R Y S V I V E A N Q T A A N Y W I					
951	960	969	978	987	996
CGT GCA CCA ATG ACC GTT GCA GGA GCC GGA ACC AAT GCA AAC TTG GAC CCC ACC					
R A P M T V A G A G T N A N L D P T					
1005	1014	1023	1032	1041	1050
AAT GTC TTT GCC GTA TTG CAC TAC GAG GGA GCG CCC AAC GCC GAA CCC ACG ACG					
N V F A V L H Y E G A P N A E P T T					
1059	1068	1077	1086	1095	1104
GAA CAA GGC AGT GCT ATC GGT ACT GCA CTC GTT GAA GAG AAC CTC CAT GCG CTC					
E Q G S A I G T A L V E E N L H A L					
1113	1122	1131	1140	1149	1158
ATC AAC CCT GGC GCT CCG GGC GGC TCC GCT CCC GCA GAC GPT TCC CTC AAT CTT					
I N P G A P G G G S A P A D V S L N L					

FIG. 4D

1167	1176	1185	1194	1203	1212
GCA ATT GGG CGC AGC ACA GGT GAT GGG ATT CTT AGG TTC ACA TTT AAT AAC ATC					
---	---	---	---	---	---
A I G R S T V D G I L R F T F N N I					
1221	1230	1239	1248	1257	1266
AAG TAC GAG GCT CCT TCG TTG CCC ACG CTC TTG AAG ATT TTG GCA AAC AAT GCG					
---	---	---	---	---	---
K Y E A P S L P T L L K I L A N N A					
1275	1284	1293	1302	1311	1320
AGC AAT GAC GCC GAT TTC ACG CCA AAT GAG CAC ACT ATC GTA TTG CCA CAC AAT					
---	---	---	---	---	---
S N D A D F T P P N E H T I V L P H N					
1329	1338	1347	1356	1365	1374
AAA GTT ATC GAG CTC AAT ATC ACC GGA GGT GCA GAC CAC CCT ATC CAT CTC CAC					
---	---	---	---	---	---
K V I E L N I T G G A D H P I H L H					
1383	1392	1401	1410	1419	1428
GGC CAT GTG TTT GAT ATC GTC AAA TCA CTC GGT GGT ACC CCG AAC TAT GTC AAC					
---	---	---	---	---	---
G H V F D I V K S L G G T P N Y V N					

FIG. 4E

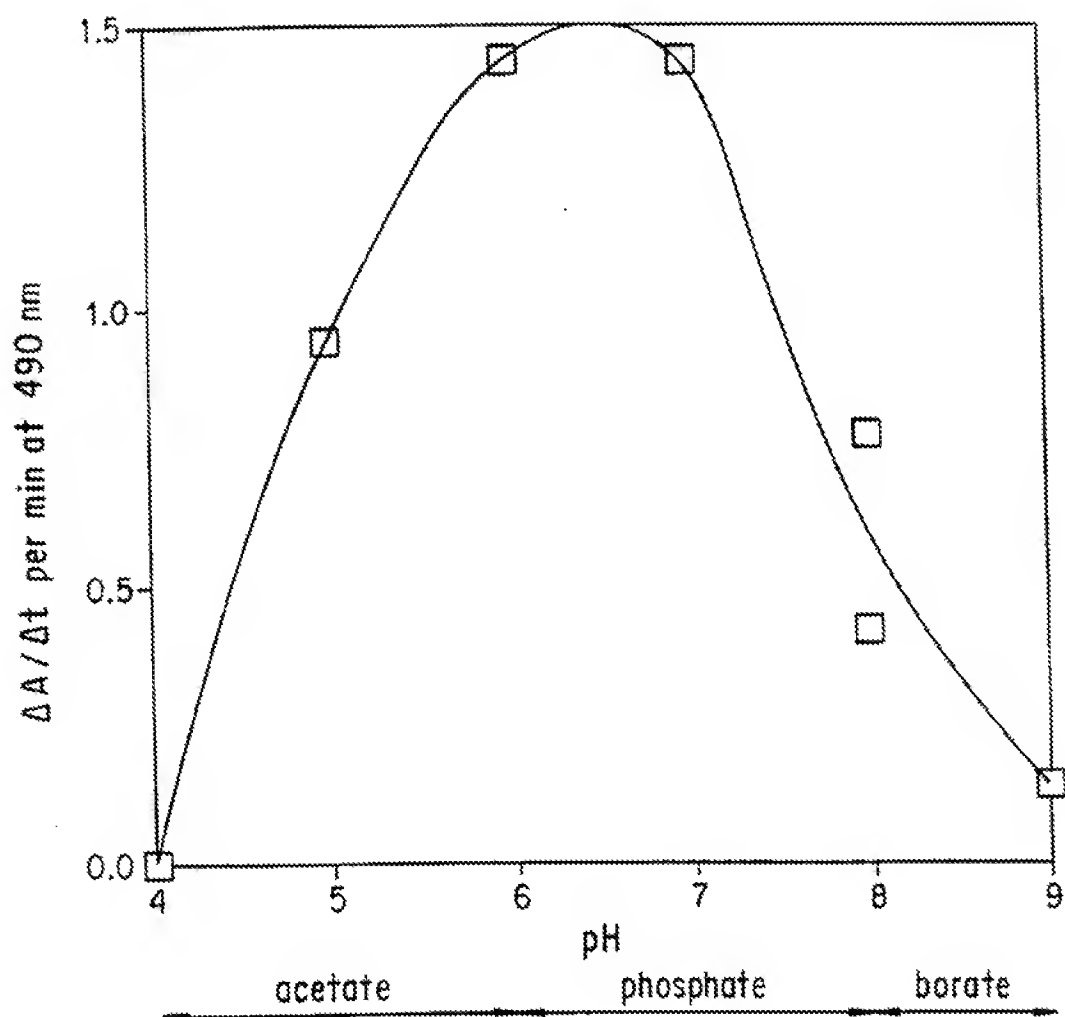


FIG. 5

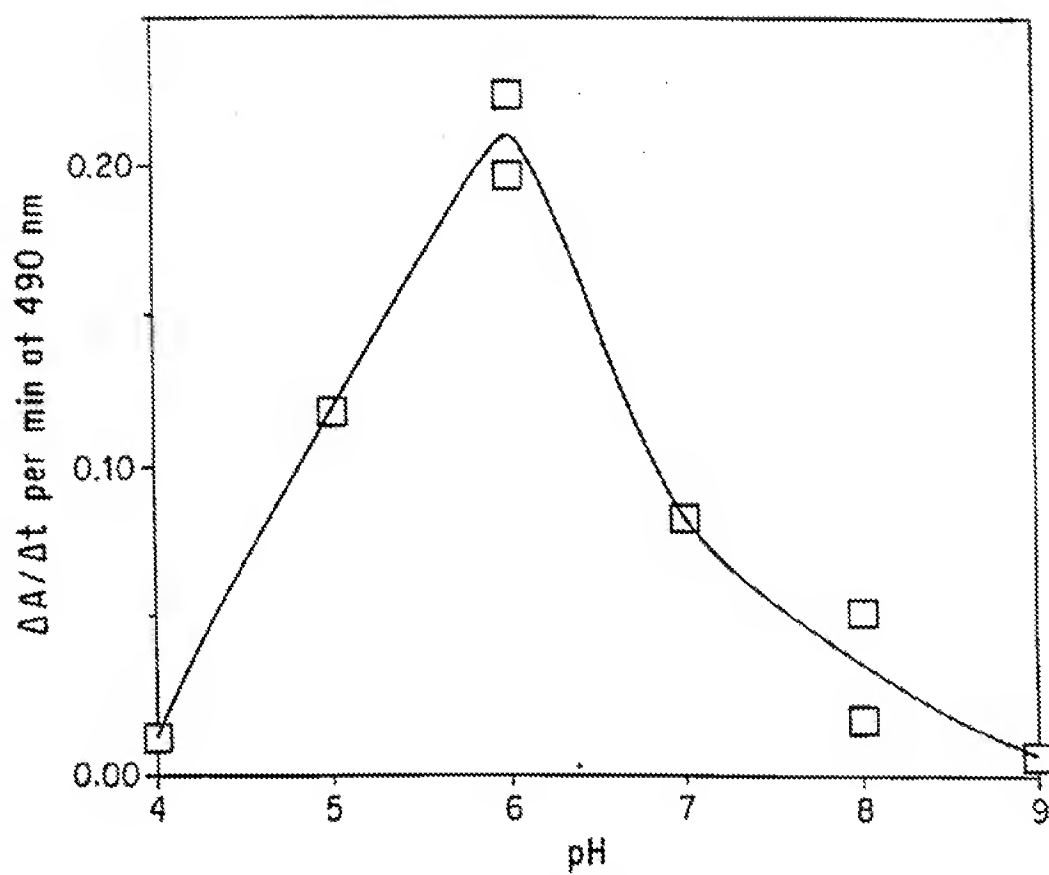


FIG. 6

INTERNATIONAL SEARCH REPORT

Int. Appl. No.

PCT/US 94/10264

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C12N15/53 C12N9/02 C12N15/80 D21C5/00 A61K7/06
 C12P7/22 C12N1/19 C09B69/10 //(C12N1/19,C12R1:66)

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C12N D21C A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	CHEMICAL ABSTRACTS, vol. 90, no. 19, 7 May 1979, Columbus, Ohio, US; abstract no. 147536w, BOLLAG J.M. ET AL. 'Characterization of an enzyme from Rhizoctonia praticola which polymerizes phenolic compounds.' page 213 ; see abstract	14,43
Y	& CAN. JOURNAL MICROBIOL., vol.25, no.2, 1979 pages 229 - 223 --- -/-	1,20-24, 39-41

☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

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 "&" document member of the same patent family

Date of the actual completion of the international search

24 January 1995

Date of mailing of the international search report

23.02.95

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 Fax (+ 31-70) 340-3016

Authorized officer

Delanghe, L

INTERNATIONAL SEARCH REPORT

Int. Appl. No.

PCT/US 94/10264

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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Information on patent family members

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PCT/US 94/10264

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EP-A-0504005	16-09-92	FR-A- 2673534 JP-A- 6172145	11-09-92 21-06-94